

NOVEL INTEGRIN α SUBUNIT AND USES THEREOFRelated Application Information

5 This application claims priority from U.S. application serial number 09/322,790, filed May 28, 1999, and from U.S. application serial number _____ (not yet available), filed April 27, 2000, both of which are hereby incorporated by reference.

Background of the Invention

10 The integrins are a family of heterodimeric membrane glycoproteins expressed on diverse cell types which function as the major receptors for extracellular matrix and as cell-cell adhesion molecules. These serve as adhesion molecules and play roles in such biological processes as platelet aggregation, inflammation, immune function, wound healing, tumor metastasis, and tissue migration during embryogenesis. There is increasing evidence to implicate integrins in signalling pathways,
15 transmitting signals both into and out of cells (Pigott, Rod and Power, Christine *The Adhesion Molecule Facts Book* (1993)).

 All integrins have two non-covalently bound subunits, referred to as alpha and beta subunits. To date, there have been 16 alpha (α) and 8 beta (β) subunits identified, the combination of which have resulted in 22 integrins seen in nature. Not all combinations of α and β subunits are possible.
20 The β subunits are more closely homologous as a group than the α subunits, and exhibit anywhere from 40% to 48% amino acid homology to one another. β subunits typically have a short (about 40-50 amino acids) cytoplasmic domain.

 The α subunits are less homologous to one another, though all contain seven repeating domains in their extracellular domains that are about 24-45 amino acids in length and are spaced
25 about 20-35 amino acids apart. These repeat domains are predicted to fold into a β -propeller domain, with the last three or four repeats containing putative divalent cation binding sites. α subunits undergo proteolytic cleavage into heavy and light fragments that are disulfide linked, unless there is an I-domain present. The I-domain, or inserted domain, is about 200 amino acids in length, typically appears between the 2nd and 3rd repeat domain, contains a metal ion-dependent adhesion
30 site, and is implicated in ligand binding. The I-domain is similar to a von Willebrand factor domain.

 Integrins can bind a wide range of ligands, common types of which include fibronectin and laminin. Integrins tend to bind with low affinity, and as a result can only bind their ligands when present in large numbers at certain regions of the cell surface called focal contacts and hemidesmosomes. Both subunits are responsible for ligand binding. In addition, the presence of
35 cations is necessary for ligand binding. Binding usually occurs on the β subunit, but is modified by

metal binding to the α subunit, which may also mediate specificity of ligand binding. α subunits have 3 or 4 divalent cation binding sites at their N-terminus, usually found among the three or four repeat domains found more extracellularly.

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Summary of the Invention

This invention provides novel human and murine nucleic acid molecules which encode proteins referred to herein as A259 proteins, which are novel integrin α subunits. These proteins, fragments, derivatives, and variants thereof are collectively referred to herein as "polypeptides of the invention" or "proteins of the invention." Nucleic acid molecules encoding the polypeptides or proteins of the invention are collectively referred to as "nucleic acids of the invention."

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The A259 proteins are homologous to the α subunits of the integrin family, and in particular, to the $\alpha 10$ and $\alpha 1$ subunits of the integrin family.

The nucleic acids and polypeptides of the present invention are useful as modulating agents in regulating a variety of cellular processes. Accordingly, in one aspect, this invention provides isolated nucleic acid molecules encoding a polypeptide of the invention or a biologically active portion thereof. The present invention also provides nucleic acid molecules which are suitable for use as primers or hybridization probes for the detection of nucleic acids encoding a polypeptide of the invention.

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The invention features nucleic acid molecules which are at least 47% (or 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, or 98%) identical to the nucleotide sequence of SEQ ID NO:1, SEQ ID NO:19, the nucleotide sequence of the cDNA insert of a clone deposited with ATCC as Accession Number 207190 or 207191, or a complement thereof.

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The invention features nucleic acid molecules which are at least 56% (or 57%, 58%, 59%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, or 98%) identical to the nucleotide sequence of SEQ ID NO:2, SEQ ID NO:20, the nucleotide sequence of the cDNA insert of a clone deposited with ATCC as Accession Number 207190 or 207191, or a complement thereof.

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The invention features nucleic acid molecules which include a fragment of at least 390 (400, 500, 600, 800, 1000, 1250, 1500, 1750, 2000, 2250, 2500, 2750, 3000, 3250, 3500, 3750, 4000, 4250, 4500, 4750, 5000, 5025, or 5042) nucleotides of the nucleotide sequence of SEQ ID NO:1, SEQ ID NO:19, the nucleotide sequence of the cDNA of ATCC Accession Number 207190 or 207191, or a complement thereof.

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The invention also features nucleic acid molecules which include a nucleotide sequence encoding a protein having an amino acid sequence that is at least 44% (or 45%, 50%, 55%, 60%, 65%, 75%, 85%, 95%, or 98%) identical to the amino acid sequence of SEQ ID NO:3, SEQ ID

NO:21, the amino acid sequence encoded by the cDNA of ATCC Accession Number 207190 or 207191, or a complement thereof.

In preferred embodiments, the nucleic acid molecules have the nucleotide sequence of SEQ ID NO:1, 2, 19, 20, or the nucleotide sequence of the cDNA of ATCC Accession Number 207190 or 207191.

Also within the invention are nucleic acid molecules which encode a fragment of a polypeptide having the amino acid sequence of SEQ ID NO:3, SEQ ID NO:21, or a fragment including at least 15 (25, 30, 50, 75, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1050, 1100, 1150, 1170, 1180, or 1188) contiguous amino acids of SEQ ID NO:3, SEQ ID NO:21, or the amino acid sequence encoded by the cDNA of ATCC Accession Number 207190 or 207191.

The invention includes nucleic acid molecules which encode a naturally occurring allelic variant of a polypeptide comprising the amino acid sequence of SEQ ID NO:3, SEQ ID NO:21, or the amino acid sequence encoded by the cDNA of ATCC Accession Number 207190 or 207191, wherein the nucleic acid molecule hybridizes to a nucleic acid molecule consisting of a nucleic acid sequence encoding SEQ ID NO:3, SEQ ID NO:21, the amino acid sequence encoded by the cDNA of ATCC Accession Number 207190 or 207191, or a complement thereof under stringent conditions.

Also within the invention are isolated polypeptides or proteins having an amino acid sequence that is at least about 44%, preferably 45%, 50%, 55%, 60%, 65%, 75%, 85%, 95%, or 98% identical to the amino acid sequence of SEQ ID NO:3, SEQ ID NO:21, or the amino acid sequence encoded by the cDNA of ATCC Accession Number 207190 or 207191.

Also within the invention are isolated polypeptides or proteins which are encoded by a nucleic acid molecule having a nucleotide sequence that is at least about 56%, preferably 57%, 58%, 59%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, or 98% identical to the nucleic acid sequence encoding SEQ ID NO:3, SEQ ID NO:21, and isolated polypeptides or proteins which are encoded by a nucleic acid molecule having a nucleotide sequence which hybridizes under stringent hybridization conditions to a nucleic acid molecule having the nucleotide sequence of SEQ ID NO:1, 2, 19, or 20, a complement thereof, or the non-coding strand of the cDNA of ATCC Accession Number 207190 or 207191.

Also within the invention are polypeptides which are naturally occurring allelic variants of a polypeptide that includes the amino acid sequence of SEQ ID NO:3, or the amino acid sequence encoded by the cDNA of ATCC Accession Number 207190 or 207191, wherein the polypeptide is encoded by a nucleic acid molecule which hybridizes to a nucleic acid molecule having the sequence of SEQ ID NO:1, 2, 19, 20, or a complement thereof under stringent conditions.

The invention also features nucleic acid molecules that hybridize under stringent conditions to a nucleic acid molecule having the nucleotide sequence of SEQ ID NO:1, 2, 19, or 20, the cDNA of ATCC Accession Number 207190 or 207191, or a complement thereof. In other embodiments, the nucleic acid molecules are at least 390 (400, 500, 600, 800, 1000, 1250, 1500, 1750, 2000, 2250, 2500, 2750, 3000, 3250, 3500, 3750, 4000, 4250, 4500, 4750, 5000, 5025, or 5042) nucleotides in length and hybridize under stringent conditions to a nucleic acid molecule comprising the nucleotide sequence of SEQ ID NO:1, 2, 19, or 20, the cDNA of ATCC Accession Number 207190 or 207191, or a complement thereof.

In one embodiment, the invention provides an isolated nucleic acid molecule which is antisense to the coding strand of a nucleic acid of the invention.

Another aspect of the invention provides vectors, e.g., recombinant expression vectors, comprising a nucleic acid molecule of the invention. In another embodiment, the invention provides host cells containing such a vector or a nucleic acid molecule of the invention. The invention also provides methods for producing a polypeptide of the invention by culturing, in a suitable medium, a host cell of the invention containing a recombinant expression vector such that a polypeptide is produced.

Another aspect of this invention features isolated or recombinant proteins and polypeptides of the invention. Preferred proteins and polypeptides possess at least one biological activity possessed by the corresponding naturally-occurring human polypeptide. An activity, a biological activity, or a functional activity of a polypeptide or nucleic acid of the invention refers to an activity exerted by a protein, polypeptide or nucleic acid molecule of the invention on a responsive cell as determined *in vivo*, or *in vitro*, according to standard techniques. Such activities can be a direct activity, such as an association with or an enzymatic activity on a second protein, or an indirect activity, such as a cellular signaling activity mediated by interaction of the protein with a second protein.

A259 biological activities include, e.g., (1) the ability to form protein-protein interactions with proteins in the signaling pathway of the naturally-occurring polypeptide; (2) the ability to bind a ligand of the naturally-occurring polypeptide; (3) the ability to interact with an A259 ligand; and (4) the ability to modulate function, survival, morphology, migration, proliferation and/or differentiation of cells, e.g., of tissues in which it is expressed (e.g., osteoblasts, bone marrow, neural tissue).

A259 biological activities also include, e.g., (1) the ability to modulate, e.g., stabilize, protein-protein interactions (e.g., homophilic and/or heterophilic), and protein-ligand interactions, e.g., in receptor-ligand recognition; (2) ability to modulate signalling pathways; (3) ability to modulate cell-cell interactions, e.g., by acting as a cell-cell adhesion molecule, or by acting as a receptor for cell-cell adhesion molecules; (4) the ability to modulate interactions between cells and proteins (e.g., extracellular matrix proteins, e.g., collagens, fibrinogens, laminins, and fibronectin),

e.g., by acting as a cell surface receptor; (5) the ability to interact, e.g., noncovalently interact, with an integrin β subunit; and (6) the ability to exhibit an activity of an integrin $\alpha 10$ or an integrin $\alpha 1$ subunit.

Still other A259 biological activities include, e.g., (1) the ability to modulate, e.g., initiate, an immune response, e.g., an inflammatory response; (2) the ability to modulate, e.g., initiate wound healing, e.g., by modulating platelet aggregation or by modulating fibroblast attachment to wound sites during wound contraction; and (3) the ability to stimulate fibrogenesis.

In one embodiment, a polypeptide of the invention has an amino acid sequence sufficiently identical to an identified domain of a polypeptide of the invention. As used herein, the term "sufficiently identical" refers to a first amino acid or nucleotide sequence which contains a sufficient or minimum number of identical or equivalent (e.g., with a similar side chain) amino acid residues or nucleotides to a second amino acid or nucleotide sequence such that the first and second amino acid or nucleotide sequences have a common structural domain and/or common functional activity. For example, amino acid or nucleotide sequences which contain a common structural domain having about 60% identity, preferably 65% identity, more preferably 75%, 85%, 95%, 98% or more identity are defined herein as sufficiently identical.

In one embodiment, an A259 protein includes at least one or more of the following domains: a signal sequence, an extracellular domain, a repeat domain, an I domain, an integrin alpha repeat domain, a transmembrane domain, and a cytoplasmic domain. In yet another embodiment, an A259 protein includes an extracellular domain and one or more repeat domains, and or one or more integrin alpha repeat domains, and is a soluble protein. In still another embodiment, an A259 protein includes an extracellular domain, one or more repeat domains, a transmembrane domain, a cytoplasmic domain, and is a receptor protein.

The polypeptides of the present invention, or biologically active portions thereof, can be operably linked to a heterologous amino acid sequence to form fusion proteins. The invention further features antibodies that specifically bind a polypeptide of the invention such as monoclonal or polyclonal antibodies. In addition, the polypeptides of the invention or biologically active portions thereof can be incorporated into pharmaceutical compositions, which optionally include pharmaceutically acceptable carriers.

In another aspect, the present invention provides methods for detecting the presence of the activity or expression of a polypeptide of the invention in a biological sample by contacting the biological sample with an agent capable of detecting an indicator of activity such that the presence of activity is detected in the biological sample.

In another aspect, the invention provides methods for modulating activity of a polypeptide of the invention comprising contacting a cell with an agent that modulates (inhibits or stimulates) the

activity or expression of a polypeptide of the invention such that activity or expression in the cell is modulated. In one embodiment, the agent is an antibody that specifically binds to a polypeptide of the invention.

In another embodiment, the agent modulates expression of a polypeptide of the invention by modulating transcription, splicing, or translation of an mRNA encoding a polypeptide of the invention. In yet another embodiment, the agent is a nucleic acid molecule having a nucleotide sequence that is antisense to the coding strand of an mRNA encoding a polypeptide of the invention.

The present invention also provides methods to treat a subject having a disorder characterized by aberrant activity of a polypeptide of the invention or aberrant expression of a nucleic acid of the invention by administering an agent which is a modulator of the activity of a polypeptide of the invention or a modulator of the expression of a nucleic acid of the invention to the subject. In one embodiment, the modulator is a protein of the invention. In another embodiment, the modulator is a nucleic acid of the invention. In other embodiments, the modulator is a peptide, peptidomimetic, or other small molecule.

The present invention also provides diagnostic assays for identifying the presence or absence of a genetic lesion or mutation characterized by at least one of: (i) aberrant modification or mutation of a gene encoding a polypeptide of the invention, (ii) mis-regulation of a gene encoding a polypeptide of the invention, and (iii) aberrant post-translational modification of the invention wherein a wild-type form of the gene encodes a protein having the activity of the polypeptide of the invention.

In another aspect, the invention provides a method for identifying a compound that binds to or modulates the activity of a polypeptide of the invention. In general, such methods entail measuring a biological activity of the polypeptide in the presence and absence of a test compound and identifying those compounds which alter the activity of the polypeptide.

The invention also features methods for identifying a compound which modulates the expression of a polypeptide or nucleic acid of the invention by measuring the expression of the polypeptide or nucleic acid in the presence and absence of the compound.

Other features and advantages of the invention will be apparent from the following detailed description and claims.

Brief Description of the Drawings

Figures 1A-1G depict the cDNA sequence of human A259 (SEQ ID NO:1) and the predicted amino acid sequence of A259 (SEQ ID NO:3). The open reading frame of SEQ ID NO:1 extends from nucleotide 127 to nucleotide 3690 of SEQ ID NO:1 (SEQ ID NO:2).

Figure 2 depicts a hydropathy plot of human A259. Relatively hydrophobic regions of the protein are above the dashed horizontal line, and relatively hydrophilic regions of the protein are below the dashed horizontal line. The cysteine residues (cys) and potential N-glycosylation sites (Ngly) are indicated by short vertical lines just below the hydropathy trace. The dashed vertical line separates the signal sequence (amino acids 1 to 22 of SEQ ID NO:3; SEQ ID NO:5) on the left from the mature protein (amino acids 23 to 1188 of SEQ ID NO:3; SEQ ID NO:4) on the right. Thicker gray horizontal bars below the dashed horizontal line indicate extracellular ("out"), transmembrane ("TM"), and intracellular ("in") regions of the molecule.

Figures 3A-3G depict an alignment of the nucleotide sequence of human integrin $\alpha 10$ (SEQ ID NO:17; GenBank Accession Number AF074015) and the nucleotide sequence of human A259 (SEQ ID NO:1). The nucleotide sequences of human $\alpha 10$ and human A259 are 46.3% identical. This alignment was performed using the ALIGN alignment program with a PAM120 scoring matrix, a gap length penalty of 12, and a gap penalty of 4.

Figures 4A-4B depict an alignment of the amino acid sequence of human $\alpha 10$ integrin (SEQ ID NO:18; GenBank Accession Number AF074015) and the amino acid sequence of human A259 (SEQ ID NO:3). The amino acid sequences of human $\alpha 10$ and human A259 are 43.0% identical. This alignment was performed using the ALIGN alignment program with a PAM120 scoring matrix, a gap length penalty of 12, and a gap penalty of 4.

Figures 5A-5G depict the cDNA sequence of murine A259 (SEQ ID NO:19) and the predicted amino acid sequence of A259 (SEQ ID NO:21). The open reading frame of SEQ ID NO:19 extends from nucleotide 28 to nucleotide 3591 of SEQ ID NO:19 (SEQ ID NO:20).

Figure 6 depicts a hydropathy plot of murine A259. Relatively hydrophobic regions of the protein are above the dashed horizontal line, and relatively hydrophilic regions of the protein are below the dashed horizontal line. The cysteine residues (cys) and potential N-glycosylation sites (Ngly) are indicated by short vertical lines just below the hydropathy trace. The dashed vertical line separates the signal sequence (amino acids 1 to 22 of SEQ ID NO:21; SEQ ID NO:23) on the left from the mature protein (amino acids 23 to 1188 of SEQ ID NO:21; SEQ ID NO:22) on the right. Thicker gray horizontal bars below the dashed horizontal line indicate extracellular ("out"), transmembrane ("TM"), and intracellular ("in") regions of the molecule.

Figures 7A-7G depict an alignment of the nucleotide sequence of murine A259 (SEQ ID NO:19) and the nucleotide sequence of human A259 (SEQ ID NO:1). The nucleotide sequences of murine A259 and human A259 are 78.4% identical. This alignment was performed using the ALIGN alignment program with a PAM120 scoring matrix, a gap length penalty of 12, and a gap penalty of 4. There is also an 86.8% identity between the human and murine A259 open reading frames.

Figures 8A-8B depict an alignment of the amino acid sequence of murine A259 (SEQ ID NO:21) and the amino acid sequence of human A259 (SEQ ID NO:3). The amino acid sequences of murine A259 and human A259 are 90.3% identical. This alignment was performed using the ALIGN alignment program with a PAM120 scoring matrix, a gap length penalty of 12, and a gap penalty of 4.

Figure 9A depicts an alignment of amino acids 37-90 of human A259 (SEQ ID NO:35) with a consensus integrin alpha repeat domain derived from a hidden Markov model (SEQ ID NO:40).

Figure 9B depicts an alignment of amino acids 421-472 of human A259 (SEQ ID NO:36) with a consensus integrin alpha repeat domain derived from a hidden Markov model (SEQ ID NO:40).

Figure 9C depicts an alignment of amino acids 476-532 of human A259 (SEQ ID NO:37) with a consensus integrin alpha repeat domain derived from a hidden Markov model (SEQ ID NO:40).

Figure 9D depicts an alignment of amino acids 538-593 of human A259 (SEQ ID NO:38) with a consensus integrin alpha repeat domain derived from a hidden Markov model (SEQ ID NO:40).

Figure 9E depicts an alignment of amino acids 600-654 of human A259 (SEQ ID NO:39) with a consensus integrin alpha repeat domain derived from a hidden Markov model (SEQ ID NO:40).

Figure 10A is a graph depicting the results of an analysis of A259 expression in three normal liver clinical samples (A, B, and C) and eight liver fibrosis clinical samples (D, E, F, G, H, I, J, and K).

Figure 10B is a graph depicting the results an analysis of human A259 expression in a variety of cells: heart (A), lung (B), liver (C), passaged stellate cells (D), quiescent stellate cells (E), stellate cells (F) stellate/F BS cells (G), NHDF fibroblasts (H); TGF-treated NHDF fibroblasts (I); NHLF fibroblasts (J); and TGF-treated NHLF fibroblasts (K).

Detailed Description of the Invention

The A259 proteins and nucleic acid molecules comprise a family of molecules having certain conserved structural and functional features. As used herein, the term “family” is intended to mean two or more proteins or nucleic acid molecules having a common structural domain and having sufficient amino acid or nucleotide sequence identity as defined herein. Family members can be from either the same or different species. For example, a family can comprises two or more proteins of human origin, or can comprise one or more proteins of human origin and one or more of non-human origin. Members of the same family may also have common structural domains.

For example, A259 proteins of the invention have signal sequences. As used herein, a “signal sequence” includes a peptide of at least about 15 or 20 amino acid residues in length which occurs at the N-terminus of secretory and membrane-bound proteins and which contains at least about 70% hydrophobic amino acid residues such as alanine, leucine, isoleucine, phenylalanine, proline, tyrosine, tryptophan, or valine. In a preferred embodiment, a signal sequence contains at least about 10 to 40 amino acid residues, preferably about 15-30 amino acid residues, more preferably about 22 amino acid residues, and has at least about 60-80%, more preferably 65-75%, and more preferably at least about 70% hydrophobic residues. A signal sequence serves to direct a protein containing such a sequence to a lipid bilayer. Thus, in one embodiment, an A259 protein contains a signal sequence at about amino acids 1 to 22 of SEQ ID NO:3 (SEQ ID NO:5). The signal sequence is cleaved during processing of the mature protein.

In another example, an A259 family member also includes one or more of the following domains: (1) an extracellular domain; (2) a transmembrane domain; and (3) a cytoplasmic domain. Thus, in one embodiment, an A259 protein contains an extracellular domain at about amino acids 1 to 1141 of SEQ ID NO:3 (SEQ ID NO:6). In another embodiment, an A259 protein contains a transmembrane domain at about amino acids 1142 to 1164 of SEQ ID NO:3 (SEQ ID NO:15). In another embodiment, an A259 protein contains a cytoplasmic domain at about amino acids 1165 to 1188 of SEQ ID NO:3 (SEQ ID NO:16).

In one embodiment, the extracellular domain of A259 can include an inserted domain (“I” domain). In another embodiment, the I domain is located between integrin α -subunit repeat domains 2 and 3.

In a preferred embodiment, an A259 family member has the amino acid sequence of SEQ ID NO:3 wherein the extracellular domain is located at about amino acids 1 to 1141 (SEQ ID NO:6), the I domain (SEQ ID NO:7) is located at about amino acid positions 164 to 345, and the integrin α -subunit repeat domains on either side of the I domain (integrin α -subunit repeat domains 2 and 3) are located at amino acid positions 115 to 157 (SEQ ID NO:9) and 367 to 392 (SEQ ID NO:10), respectively.

An A259 family member can include one or more integrin α -subunit repeat domains within the extracellular domain. As used herein, a “repeat domain” refers to one or more of seven homologous protein domains that are conserved in integrin α -subunit family members, which are about 10 to 65 residues, preferably about 20 to 50 residues, more preferably about 25 to 45 amino acid residues, and most preferably about 26 to 43 residues, in length, and which are about 10 to 50, preferably about 15 to 40, more preferably about 20 to 35 amino acids apart from one another.

A repeat domain typically has one of the two following consensus sequences. The first repeat consensus sequence (type 1) is: F-G-Xaa(n)-[G or A]-A-[P or L or Q], wherein F is

phenylalanine, G is glycine, Xaa is any amino acid, n is about 10 to 30 amino acid residues, preferably about 15 to 20 amino acid residues, more preferably about 16 to 19 amino acid residues, A is alanine, P is proline, L is leucine, and Q is glutamine. The second repeat consensus sequence (type 2) is: [G or S]-[F or Y]-Xaa(n)-[G or V or L]-[A or M]-[P or Y], wherein G is glycine, S is serine, F is phenylalanine, Y is tyrosine, Xaa is any amino acid, n is about 5 to 40 amino acid residues, preferably about 10 to 35 amino acid residues, more preferably about 14 to 33 amino acid residues, V is valine, A is alanine, M is methionine, and P is proline.

In one embodiment, an A259 family member includes one or more repeat domains having an amino acid sequence that is at least about 55%, preferably at least about 65%, more preferably at least about 75%, yet more preferably at least about 85%, and most preferably at least about 95% identical to amino acids 39 to 74, and/or 115 to 157, and/or 367 to 392, and/or 421 to 455, and/or 478 to 516, and/or 540 to 575, and/or 602 to 640 of SEQ ID NO:3, which are the repeat domains of A259 (these repeat domains are also represented as SEQ ID NO:8, 9, 10, 11, 12, 13, and 14, respectively). In another embodiment, an A259 family member includes one or more repeat domains having an amino acid sequence that is at least about 55%, preferably at least about 65%, more preferably at least about 75%, yet more preferably at least about 85%, and most preferably at least about 95% identical to amino acids 39 to 74, and/or 115 to 157, and/or 367 to 392, and/or 421 to 455, and/or 478 to 516, and/or 540 to 575, and/or 602 to 640 of SEQ ID NO:3, which are the repeat domains of A259 (these repeat domains are also represented as SEQ ID NO:8, 9, 10, 11, 12, 13, and 14, respectively), and has one or more repeat consensus sequences described herein. In another embodiment, an A259 family member includes one or more repeat domains having an amino acid sequence that is at least about 55%, preferably at least about 65%, more preferably at least about 75%, yet more preferably at least about 85%, and most preferably at least about 95% identical to amino acids 39 to 74, and/or 115 to 157, and/or 367 to 392, and/or 421 to 455, and/or 478 to 516, and/or 540 to 575, and/or 602 to 640 of SEQ ID NO:3, which are the repeat domains of A259 (these repeat domains are also represented as SEQ ID NO:8, 9, 10, 11, 12, 13, and 14, respectively), has one or more I consensus sequences described herein, and has at least one A259 biological activity as described herein.

An A259 family member can also include an inserted, or I domain (also called von Willebrand factor type A domain). As used herein, an "I domain" refers to a domain that appears in only some of the integrin α subunits, e.g., $\alpha 1$, $\alpha 2$, αM , and αX , and that is "inserted" between the second and third integrin α -subunit repeat domains. I domains prevent the occurrence of proteolytic cleavage that separates integrin α subunits into heavy and light fragments that are disulfide linked. An I domain includes about 100 to 300 amino acid residues, preferably about 150 to 200 amino acid residues, more preferably about 160 to 190 amino acid residues, and most preferably about 182 amino acid residues.

An I domain typically has the following consensus sequence: D-G-S-Xaa-S-Xaa(n1)-F-Xaa(n2)-Q-Y, wherein D is aspartic acid, G is glycine, S is serine, Xaa is any amino acid, n1 is about 5 to 15, preferably about 8 to 12, more preferably about 9 to 11 amino acid residues, F is phenylalanine, n2 is about 15 to 30, preferably 18 to 28, more preferably about 20 to 26 amino acid residues, Q is glutamine, and Y is tyrosine.

In one embodiment, an A259 family member includes an I domain having an amino acid sequence that is at least about 55%, preferably at least about 65%, more preferably at least about 75%, yet more preferably at least about 85%, and most preferably at least about 95% identical to amino acids 164 to 345 of SEQ ID NO:3, which is the I domain of A259 (this I domain is also represented as SEQ ID NO:7). In another embodiment, an A259 family member includes an I domain having an amino acid sequence that is at least about 55%, preferably at least about 65%, more preferably at least about 75%, yet more preferably at least about 85%, and most preferably at least about 95% identical to amino acids 164 to 345 of SEQ ID NO:3, which is the I domain of A259 (this I domain is also represented as SEQ ID NO:7), and an I domain consensus sequence described herein. In another embodiment, an A259 family member includes an I domain having an amino acid sequence that is at least about 55%, preferably at least about 65%, more preferably at least about 75%, yet more preferably at least about 85%, and most preferably at least about 95% identical to amino acids 164 to 345 of SEQ ID NO:3, which is the I domain of A259 (this I domain is also represented as SEQ ID NO:7). In another embodiment, an A259 family member includes an I domain having an amino acid sequence that is at least about 55%, preferably at least about 65%, more preferably at least about 75%, yet more preferably at least about 85%, and most preferably at least about 95% identical to amino acids 164 to 345 of SEQ ID NO:3, which is the I domain of A259 (this I domain is also represented as SEQ ID NO:7), has one or more I domain consensus sequences described herein, and has at least one A259 biological activity as described herein.

An A259 family member can also include a cytoplasmic domain. The cytoplasmic domain typically includes about 10 to 40, preferably about 20 to 30, more preferably about 22 to 28, still more preferably about 22 to 26 amino acid residues in length. The A259 cytoplasmic domain typically has the following consensus sequence: [F or Y or W or S]-[R or K]-Xaa(n1)-G-F-F-Xaa(n2)-R, wherein F is phenylalanine, Y is tyrosine, W is tryptophan, S is serine, R is arginine, K is lysine, Xaa is any amino acid, n1 and n2 are 0 to 1, and G is glycine.

In a preferred embodiment, an A259 family member has the amino acid sequence of SEQ ID NO:3 wherein the cytoplasmic domain is located at about amino acids 1165 to 1188 (SEQ ID NO:16) and the cytoplasmic domain consensus sequence is located from about amino acid 1165 to 1170.

Various features of human and murine A259 are summarized below.

Human A259

A cDNA encoding human A259 was identified by analyzing the sequences of clones present in an LPS stimulated human primary osteoblast cDNA library. This analysis led to the identification of a clone, Atho002i17, encoding full-length human A259. The human A259 cDNA of this clone is 5042 nucleotides long (Figures 1A-1D; SEQ ID NO:1). The open reading frame of this cDNA, nucleotides 127 to 3690 of SEQ ID NO:1 (SEQ ID NO:2), encodes a 1188 amino acid receptor protein (Figures 1A-1D; SEQ ID NO:3).

The signal peptide prediction program SIGNALP (Nielsen et al. (1997) *Protein Engineering* 10:1-6) predicted that human A259 includes a 22 amino acid signal peptide (amino acid 1 to about amino acid 22 of SEQ ID NO:3; SEQ ID NO:5) preceding the mature A259 protein (corresponding to about amino acid 23 to amino acid 1188 of SEQ ID NO:3)(SEQ ID NO:4). The human A259 protein molecular weight is 133.5 kDa prior to the cleavage of the signal peptide, 131.1 kDa after cleavage of the signal peptide.

Human A259 includes a extracellular domain (about amino acids 1 to 1141 of SEQ ID NO:3; SEQ ID NO:6), an I domain (also called a Von Willebrand Factor type A domain; about amino acids 164 to 345 of SEQ ID NO:3; SEQ 7), seven repeat domains (about amino acids 39 to 74, 115 to 157, 367 to 392, 421 to 455, 478 to 516, 540 to 575, and 602 to 640 of SEQ ID NO:3 (SEQ ID NO:8, 9, 10, 11, 12, 13, and 14, respectively)), a transmembrane domain (about amino acids 1142 to 1164 of SEQ ID NO:3; SEQ ID NO:15), and a cytoplasmic domain (about amino acids 1165 to 1188 of SEQ ID NO:3; SEQ ID NO:16).

A slightly different model of the integrin alpha repeat domain identifies five integrin alpha repeat domains within human A259 (about amino acids 37-90, 421-472, 476-532, 538-593, and 600-654 of SEQ ID NO:3 (SEQ ID NOS:35, 36, 37, 38, and 39, respectively)). These domains were identified by homology searching using consensus domains derived from hidden Markov models (HMMs). HMMs can be used to do multiple sequence alignment and very sensitive database searching, using statistical descriptions of a sequence family's consensus. For more information on HMM searches, see, e.g., <http://hmmer.wustl.edu>. Figure 9A depicts an alignment of amino acids 37-90 of human A259 (SEQ ID NO:35) with a consensus integrin alpha repeat domain derived from a hidden Markov model (SEQ ID NO:40). Figure 9B depicts an alignment of amino acids 421-472 of human A259 (SEQ ID NO:36) with a consensus integrin alpha repeat domain derived from a hidden Markov model (SEQ ID NO:40). Figure 9C depicts an alignment of amino acids 476-532 of human A259 (SEQ ID NO:37) with a consensus integrin alpha repeat domain derived from a hidden Markov model (SEQ ID NO:40). Figure 9D depicts an alignment of amino acids 538-593 of human A259 (SEQ ID NO:38) with a consensus integrin alpha repeat domain derived from a hidden Markov model (SEQ ID NO:40). Figure 9E depicts an alignment of amino acids 600-654 of human A259

(SEQ ID NO:39) with a consensus integrin alpha repeat domain derived from a hidden Markov model (SEQ ID NO:40).

5 A predicted N-glycosylation site having the sequence NCTK is found from amino acids 82 to 85 of SEQ ID NO:3. A second predicted N-glycosylation site having the sequence NVSE is found from amino acids 95 to 98. A third predicted N-glycosylation site having the sequence NVTR is found from amino acids 291 to 294. A fourth predicted N-glycosylation site having the sequence NVTD is found from amino acids 331 to 334. A fifth predicted N-glycosylation site having the sequence NETS is found from amino acids 358 to 361. A sixth predicted N-glycosylation site having the sequence NHTG is found from amino acids 449 to 452. A seventh predicted N-glycosylation site
10 having the sequence NRS� is found from amino acids 462 to 465. An eighth predicted N-glycosylation site having the sequence NGTL is found from amino acids 528 to 531. A ninth predicted N-glycosylation site having the sequence NASL is found from amino acids 642 to 645. A tenth predicted N-glycosylation site having the sequence NATM is found from amino acids 694 to 697. An eleventh predicted N-glycosylation site having the sequence NISQ is found from amino
15 acids 857 to 860. A twelfth predicted N-glycosylation site having the sequence NVSY is found from amino acids 894 to 897. A thirteenth predicted N-glycosylation site having the sequence NSSL is found from amino acids 973 to 976. A fourteenth predicted N-glycosylation site having the sequence NTSC is found from amino acids 1031 to 1034. A fifteenth predicted N-glycosylation site having the sequence NSTE is found from amino acids 1039 to 1042. A sixteenth predicted N-glycosylation site
20 having the sequence NHSN is found from amino acids 1059 to 1062. A cAMP and cGMP-dependent protein kinase phosphorylation site having the sequence RRYT is found from amino acids 700 to 703.

A predicted protein kinase C phosphorylation site having the sequence TRK is found from amino acids 27 to 29 of SEQ ID NO:3. A second predicted protein kinase C phosphorylation site
25 having the sequence SER is found from amino acids 97 to 99. A third predicted protein kinase C phosphorylation site having the sequence SVK is found from amino acids 221 to 223. A fourth predicted protein kinase C phosphorylation site having the sequence SER is found from amino acids 287 to 289. A fifth predicted protein kinase C phosphorylation site having the sequence TNK is found from amino acids 355 to 357. A sixth predicted protein kinase C phosphorylation site having
30 the sequence SSR is found from amino acids 434 to 436. A seventh predicted protein kinase C phosphorylation site having the sequence TĠK is found from amino acids 451 to 453. An eighth predicted protein kinase C phosphorylation site having the sequence TLK is found from amino acids 530 to 532. A ninth predicted protein kinase C phosphorylation site having the sequence SVR is found from amino acids 548 to 550. A tenth predicted protein kinase C phosphorylation site having
35 the sequence TPK is found from amino acids 587 to 589. An eleventh predicted protein kinase C

phosphorylation site having the sequence SGR is found from amino acids 662 to 664. A twelfth predicted protein kinase C phosphorylation site having the sequence TPR is found from amino acids 703 to 705. A thirteenth predicted protein kinase C phosphorylation site having the sequence TNR is found from amino acids 716 to 718. A fourteenth predicted protein kinase C phosphorylation site having the sequence TLR is found from amino acids 770 to 772. A fifteenth predicted protein kinase C phosphorylation site having the sequence STR is found from amino acids 833 to 835. A sixteenth predicted protein kinase C phosphorylation site having the sequence STK is found from amino acids 938 to 940. A seventeenth predicted protein kinase C phosphorylation site having the sequence SLK is found from amino acids 1092 to 1094. An eighteenth predicted protein kinase C phosphorylation site having the sequence SMK is found from amino acids 1100 to 1102. A nineteenth predicted protein kinase C phosphorylation site having the sequence SAR is found from amino acids 1171 to 1173. A twentieth predicted protein kinase C phosphorylation site having the sequence TPK is found from amino acids 1183 to 1185.

A predicted casein kinase II phosphorylation site having the sequence TYMD is found from amino acids 161 to 164 of SEQ ID NO:3. A second predicted casein kinase II phosphorylation site having the sequence SVKD is found from amino acids 221 to 224. A third predicted casein kinase II phosphorylation site having the sequence SHIE is found from amino acids 230 to 233. A fourth predicted casein kinase II phosphorylation site having the sequence TDGE is found from amino acids 270 to 273. A fifth predicted casein kinase II phosphorylation site having the sequence SERD is found from amino acids 287 to 290. A sixth predicted casein kinase II phosphorylation site having the sequence SDPD is found from amino acids 322 to 325. A seventh predicted casein kinase II phosphorylation site having the sequence TSVD is found from amino acids 485 to 488. An eighth predicted casein kinase II phosphorylation site having the sequence TLKD is found from amino acids 530 to 533. A ninth predicted casein kinase II phosphorylation site having the sequence SVRD is found from amino acids 548 to 551. A tenth predicted casein kinase II phosphorylation site having the sequence SYND is found from amino acids 556 to 559. An eleventh predicted casein kinase II phosphorylation site having the sequence TASE is found from amino acids 593 to 596. A twelfth predicted casein kinase II phosphorylation site having the sequence SGRD is found from amino acids 662 to 665. A thirteenth predicted casein kinase II phosphorylation site having the sequence TMDE is found from amino acids 696 to 669. A fourteenth predicted casein kinase II phosphorylation site having the sequence SGQE is found from amino acids 724 to 727. A fifteenth predicted casein kinase II phosphorylation site having the sequence SLED is found from amino acids 753 to 756. A sixteenth predicted casein kinase II phosphorylation site having the sequence TAME is found from amino acids 801 to 804. A seventeenth predicted casein kinase II phosphorylation site having the sequence STKE is found from amino acids 938 to 941. An eighteenth predicted casein kinase II

phosphorylation site having the sequence SHYE is found from amino acids 966 to 969. A nineteenth predicted casein kinase II phosphorylation site having the sequence SSLE is found from amino acids 974 to 977. A twentieth predicted casein kinase II phosphorylation site having the sequence TPVE is found from amino acids 1046 to 1049. A twenty-first predicted casein kinase II phosphorylation site having the sequence SNSD is found from amino acids 1061 to 1064. A twenty-second predicted casein kinase II phosphorylation site having the sequence SKQE is found from amino acids 1133 to 1136.

A predicted tyrosine kinase phosphorylation site having the sequence KPAQDCSAY is found from amino acids 812 to 820.

An predicted N-myristoylation site having the sequence GLVVAW is found from amino acids 6 to 11 of SEQ ID NO:3. A second predicted N-myristoylation site having the sequence GSRTAF is found from amino acids 35 to 40. A third predicted N-myristoylation site having the sequence GLSLAT is found from amino acids 106 to 111. A fourth predicted N-myristoylation site having the sequence GGTETR is found from amino acids 236 to 241. A fifth predicted N-myristoylation site having the sequence GIEFAR is found from amino acids 245 to 250. A sixth predicted N-myristoylation site having the sequence GGRKGA is found from amino acids 257 to 262. A seventh predicted N-myristoylation site having the sequence GTNKNE is found from amino acids 354 to 359. An eighth predicted N-myristoylation site having the sequence GLEMSQ is found from amino acids 363 to 368. A ninth predicted N-myristoylation site having the sequence GVLLGA is found from amino acids 379 to 384. A tenth predicted N-myristoylation site having the sequence GAYLGY is found from amino acids 422 to 427. An eleventh predicted N-myristoylation site having the sequence GQQIGS is found from amino acids 473 to 478. A twelfth predicted N-myristoylation site having the sequence GSEITS is found from amino acids 481 to 486. A thirteenth predicted N-myristoylation site having the sequence GSSIAS is found from amino acids 543 to 548. A fourteenth predicted N-myristoylation site having the sequence GALGNA is found from amino acids 625 to 630. A fifteenth predicted N-myristoylation site having the sequence GIRYNA is found from amino acids 690 to 695. A sixteenth predicted N-myristoylation site having the sequence GQELCE is found from amino acids 725 to 730. A seventeenth predicted N-myristoylation site having the sequence GSIECV is found from amino acids 877 to 882. An eighteenth predicted N-myristoylation site having the sequence GSDSNE is found from amino acids 930 to 935. A nineteenth predicted N-myristoylation site having the sequence GSTLGG is found from amino acids 1147 to 1152.

A predicted amidation site having the sequence GGRK is found from amino acids 257 to 260 of SEQ ID NO:3. An immunoglobulin and major histocompatibility complex proteins signature having the sequence YKCPVIH is found from amino acids 74 to 80.

Human A259 is homologous to human integrin $\alpha 10$ (SEQ ID NO:17; GenBank Accession Number AF074015), a $\beta 1$ -associated collagen binding integrin (Camper, Hellman, and Lundgren-Akerlund (1998) *Journal of Biol. Chem.* 273, 32:20383-20389). Figures 3A-3H depict an alignment of the nucleotide sequence of human integrin $\alpha 10$ (SEQ ID NO:17; GenBank Accession Number AF074015) with the nucleotide sequence of human A259 (SEQ ID NO:1). Figures 4A-4B depict an alignment of the amino acid sequence of human $\alpha 10$ integrin (SEQ ID NO:18; GenBank Accession Number AF074015) with the amino acid sequence of human A259 (SEQ ID NO:3). As shown in Figures 4A-4B, the human A259 signal sequence is represented by amino acids 1 to 22 (and encoded by nucleotides 127 to 192 of SEQ ID NO:1). The human $\alpha 10$ integrin signal sequence is represented by amino acids 1 to 22 (and encoded by nucleotides 22 to 87 of SEQ ID NO:17). The human A259 extracellular domain sequence (SEQ ID NO:6) is represented by amino acids 1 to 1141 (and encoded by nucleotides 127 to 3549 of SEQ ID NO:1), and the human $\alpha 10$ extracellular domain sequence is represented by amino acids 1 to 1098 (and encoded by nucleotides 22 to 3381 of SEQ ID NO:17). The human A259 I domain (SEQ ID NO:7) is represented by amino acids 164 to 345 (and encoded by nucleotides 616 to 1161 of SEQ ID NO:1), and the human $\alpha 10$ I domain is represented by amino acids 140 to 337 (and encoded by nucleotides 505 to 1098 of SEQ ID NO:17). The human A259 transmembrane domain (SEQ ID NO:15) is represented by amino acids 1142 to 1164 (and encoded by nucleotides 3550 to 3618 of SEQ ID NO:1), and the human $\alpha 10$ transmembrane domain is represented by amino acids 1099 to 1123 (and encoded by nucleotides 3382 to 3456 of SEQ ID NO:17). The human A259 cytoplasmic domain (SEQ ID NO:16) is represented by amino acids 1165 to 1188 (and encoded by nucleotides 3619 to 3690 of SEQ ID NO:1), and the human $\alpha 10$ cytoplasmic domain is represented by amino acids 1124 to 1145 (and encoded by nucleotides 3457 to 3522 of SEQ ID NO:17).

Figures 3A-3H and 4A-4B show that there is an overall 46.3% identity between the full length human A259 nucleic acid molecule and the full length human $\alpha 10$ molecule, and a 43.0% identity between the A259 amino acid sequence and the human $\alpha 10$ amino acid sequence. There is also a 55.1% identity between the human A259 and the human $\alpha 10$ open reading frames. There is also an overall 37.6% amino acid identity, and a 41.5% full length nucleic acid identity, between the full length human A259 nucleic acid molecule and the full length human $\alpha 1$ (VLA-1 (very late antigen-1) molecule).

Human A259 is homologous to murine A259. Figures 7A-7G depict an alignment of the nucleotide sequence of murine A259 (SEQ ID NO:19) with the nucleotide sequence of human A259 (SEQ ID NO:1). Figures 8A-8B depict an alignment of the amino acid sequence of murine A259 (SEQ ID NO:21) with the amino acid sequence of human A259 (SEQ ID NO:3). As shown in Figures 8A-8B, the human A259 signal sequence is represented by amino acids 1 to 22 (and encoded

by nucleotides 127 to 192 of SEQ ID NO:1). The murine A259 signal sequence is represented by amino acids 1 to 22 (and encoded by nucleotides 28 to 93 of SEQ ID NO:19). The human A259 extracellular domain sequence (SEQ ID NO:6) is represented by amino acids 1 to 1141 (and encoded by nucleotides 127 to 3549 of SEQ ID NO:1), and the murine A259 extracellular domain sequence is represented by amino acids 1 to 1141 (and encoded by nucleotides 28 to 3450 of SEQ ID NO:19). The human A259 I domain (SEQ ID NO:7) is represented by amino acids 164 to 345 (and encoded by nucleotides 616 to 1161 of SEQ ID NO:1), and the murine A259 I domain is represented by amino acids 164 to 345 (and encoded by nucleotides 517 to 1062 of SEQ ID NO:19). The human A259 repeat domains are represented by amino acids 39 to 74, 115 to 157, 367 to 392, 421 to 455, 478 to 516, 540 to 575, and 602 to 640 (and encoded by nucleotides 244 to 348, 469 to 597, 1225 to 1302, 1387 to 1491, 1558 to 1674, 1744 to 1851, and 1930 to 2046, respectively, of SEQ ID NO:1), and the murine A259 repeat domains are represented by amino acids 39 to 74, 115 to 157, 367 to 392, 421 to 455, 478 to 516, 540 to 575, and 602 to 640 (and encoded by nucleotides 145 to 249, 370 to 498, 1126 to 1203, 1288 to 1392, 1459 to 1575, 1645 to 1752, and 1831 to 1947, respectively, of SEQ ID NO:19). The human A259 transmembrane domain (SEQ ID NO:15) is represented by amino acids 1142 to 1164 (and encoded by nucleotides 3550 to 3618 of SEQ ID NO:1), and the murine A259 transmembrane domain is represented by amino acids 1142 to 1164 (and encoded by nucleotides 3283 to 3357 of SEQ ID NO:19). The human A259 cytoplasmic domain (SEQ ID NO:16) is represented by amino acids 1165 to 1188 (and encoded by nucleotides 3619 to 3690 of SEQ ID NO:1), and the murine A259 cytoplasmic domain is represented by amino acids 1165 to 1188 (and encoded by nucleotides 3358 to 3423 of SEQ ID NO:19).

Figures 7A-7G and 8A-8B show that there is an overall 78.4% identity between the full length human A259 nucleic acid molecule and the full length murine A259 molecule, and a 90.3% identity between the A259 amino acid sequence and the murine A259 amino acid sequence. There is also an 86.8% identity between the human and murine A259 open reading frames.

Clone Human I2, which encodes human A259, was deposited with the American Type Culture Collection (10801 University Boulevard, Manassas, VA 20110-2209) on April 2, 1999 and assigned Accession Number 207190. This deposit will be maintained under the terms of the Budapest Treaty on the International Recognition of the Deposit of Microorganisms for the Purposes of Patent Procedure. This deposit was made merely as a convenience for those of skill in the art and is not an admission that a deposit is required under 35 U.S.C. §112.

Figure 2 depicts a hydropathy plot of human A259. Relatively hydrophobic regions of the protein are shown above the horizontal line, and relatively hydrophilic regions of the protein are below the horizontal line. The cysteine residues (cys) and predicted N-glycosylation sites are indicated by short vertical lines just below the hydropathy trace. The dashed vertical line separates

the signal sequence (amino acids 1 to 22 of SEQ ID NO:3; SEQ ID NO:5) on the left from the mature protein (amino acids 23 to 1188 of SEQ ID NO:3; SEQ ID NO:4) on the right. The A259 transmembrane domain is indicated by the section of the plot under which the number 6.4 can be seen, which represents a score assigned to the predicted transmembrane domain. The extracellular domain (SEQ ID NO:6) and cytoplasmic domain (SEQ ID NO:16) are similarly indicated by gray horizontal bars, labeled as Aout≡ and Ain≡, respectively.

An electronic survey of proprietary databases revealed that A259 is found in abundantly in human and mouse osteoblast cDNA libraries and also in human bone marrow and neuronal cDNA libraries. In addition, in a Northern blot containing RNA from human tissues, A259 was expressed as two messages, a 6kb and a 4kb message, in RNA from human osteoblasts. The sequence used to probe this human Northern blot contained human A259 C-terminal and 3' untranslated regions. The expression level in human osteoblasts did not change when stimulated with TNF- α .

The expression of human A259 was measured in various clinical liver samples taken from subjects not suffering from liver fibrosis and from patients suffering from liver fibrosis using TaqMan quantitative PCR. The results of this analysis are depicted in Figure 10A. Human A259 was expressed at a lower level in the livers of patients not suffering from liver fibrosis (A, B, and C; relative expression 0.13, 0.29, and .09, respectively) than in the livers of patients suffering from liver fibrosis (D, E, F, G, H, I, J, and K; relative expression 1.59, 0.50, 0.55, 1.56, 0.62, 0.93, 3.54, and 0.61, respectively).

The expression of human A259 was measured in various cells using TaqMan quantitative PCR. The results of this analysis are depicted in Figure 10B. The cells tested were: heart (A; relative expression 1.69), lung (B; relative expression 0.41), liver (C; relative expression 0.10), passaged stellate cells (D; relative expression 36.15), quiescent stellate cells (E; relative expression 0.15), stellate cells (F; relative expression 18.71); stellate/F BS cells (G; relative expression 31.03), NHDF fibroblasts (H; relative expression 0.71); TGF-treated NHDF fibroblasts (I; relative expression 12.69); NHLF fibroblasts (J; relative expression 4.81); and TGF-treated NHLF fibroblasts (K; relative expression 66.06).

Human A259 maps to human chromosome location hu15q22. The flanking markers are AFM094YC1 and NIB1778. The nearby known disease loci include: CLN6 (neuronal ceroid-lipofuscinosis) and BBS4 (Bardet-Biedl Syndrome 4). The nearby known genes include: HDC (histidine decarboxylase), LIPC (hepatic lipase), PKM2 (pyruvate kinase 3), CLN6 (neuronal ceroid-lipofuscinosis), BBS4 (Bardet-Biedl Syndrome 4), and NMB (neuromedin B). The murine syntenic chromosomal location is mo9. The known nearby loci include: rc (rough coat), ash (ashen), flail (flailer), pml (promyelotic leukemia), and sty (small thymus). The known nearby genes include: hpl (hepatitic lipase).

Murine A259

A cDNA encoding human A259 was identified by analyzing the sequences of clones present in a murine bone marrow cDNA library. This analysis led to the identification of a clone, AtmMa113d1, encoding full-length murine A259. The murine A259 cDNA of this clone is 4858 nucleotides long (Figures 4A-4D; SEQ ID NO:19). The open reading frame of this cDNA, nucleotides 28 to 3591 of SEQ ID NO:19 (SEQ ID NO:20), encodes a 1188 amino acid receptor protein (Figures 4A-4D; SEQ ID NO:21).

The signal peptide prediction program SIGNALP (Nielsen et al. (1997) Protein Engineering 10:1-6) predicted that murine A259 includes a 22 amino acid signal peptide (amino acid 1 to about amino acid 22 of SEQ ID NO:21)(SEQ ID NO:23) preceding the mature A259 protein (corresponding to about amino acid 23 to amino acid 1188 of SEQ ID NO:21)(SEQ ID NO:22). The mouse A259 protein molecular weight is 133.1 kDa prior to the cleavage of the signal peptide, 130.5 kDa after cleavage of the signal peptide.

Murine A259 includes a extracellular domain (about amino acids 1 to 1141 of SEQ ID NO:21; SEQ ID NO:24), an I domain (about amino acids 164 to 345 of SEQ ID NO:21; SEQ 25), seven repeat domains (about amino acids 39 to 74, 115 to 157, 367 to 392, 421 to 455, 478 to 516, 540 to 575, and 602 to 640 of SEQ ID NO:21 (SEQ ID NO:26, 27, 28, 29, 30, 31, and 32, respectively)), a transmembrane domain (about amino acids 1142 to 1164 of SEQ ID NO:21; SEQ ID NO:33), and a cytoplasmic domain (about amino acids 1165 to 1188 of SEQ ID NO:21; SEQ ID NO:34).

A predicted N-glycosylation site having the sequence NCTK is found from amino acids 82 to 85 of SEQ ID NO:21. A second predicted N-glycosylation site having the sequence NVSE is found from amino acids 95 to 98. A third predicted N-glycosylation site having the sequence NVTR is found from amino acids 291 to 294. A fourth predicted N-glycosylation site having the sequence NVTD is found from amino acids 331 to 334. A fifth predicted N-glycosylation site having the sequence NETS is found from amino acids 358 to 361. A sixth predicted N-glycosylation site having the sequence NHTG is found from amino acids 449 to 452. A seventh predicted N-glycosylation site having the sequence NRSL is found from amino acids 462 to 465. An eighth predicted N-glycosylation site having the sequence NGTL is found from amino acids 528 to 531. A ninth predicted N-glycosylation site having the sequence NASL is found from amino acids 642 to 645. A tenth predicted N-glycosylation site having the sequence NATM is found from amino acids 694 to 697. An eleventh predicted N-glycosylation site having the sequence NISQ is found from amino acids 857 to 860. A twelfth predicted N-glycosylation site having the sequence NVSY is found from amino acids 894 to 897. A thirteenth predicted N-glycosylation site having the sequence NSSL is

found from amino acids 973 to 976. A fourteenth predicted N-glycosylation site having the sequence NTSC is found from amino acids 1031 to 1034. A fifteenth predicted N-glycosylation site having the sequence NSTE is found from amino acids 1039 to 1042. A sixteenth predicted N-glycosylation site having the sequence NHSN is found from amino acids 1059 to 1062.

5 A predicted protein kinase C phosphorylation site having the sequence SGK is found from amino acids 51 to 53 of SEQ ID NO:21. A second predicted protein kinase C phosphorylation site having the sequence SER is found from amino acids 97 to 99. A third predicted protein kinase C phosphorylation site having the sequence SVK is found from amino acids 221 to 223. A fourth predicted protein kinase C phosphorylation site having the sequence SEK is found from amino acids
10 287 to 289. A fifth predicted protein kinase C phosphorylation site having the sequence TNK is found from amino acids 355 to 357. A sixth predicted protein kinase C phosphorylation site having the sequence SSR is found from amino acids 434 to 436. A seventh predicted protein kinase C phosphorylation site having the sequence TGK is found from amino acids 451 to 453. An eighth predicted protein kinase C phosphorylation site having the sequence TLK is found from amino acids
15 530 to 532. A ninth predicted protein kinase C phosphorylation site having the sequence SHR is found from amino acids 569 to 571. A tenth predicted protein kinase C phosphorylation site having the sequence TNR is found from amino acids 716 to 718. An eleventh predicted protein kinase C phosphorylation site having the sequence TLR is found from amino acids 770 to 772. A twelfth predicted protein kinase C phosphorylation site having the sequence STR is found from amino acids
20 833 to 835. A thirteenth predicted protein kinase C phosphorylation site having the sequence SLK is found from amino acids 1092 to 1094. An fourteenth predicted protein kinase C phosphorylation site having the sequence SLK is found from amino acids 1100 to 1102. A fifteenth predicted protein kinase C phosphorylation site having the sequence SAK is found from amino acids 1171 to 1173.

A predicted casein kinase II phosphorylation site having the sequence TYMD is found from
25 amino acids 161 to 164 of SEQ ID NO:21. A second predicted casein kinase II phosphorylation site having the sequence SVKD is found from amino acids 221 to 224. A third predicted casein kinase II phosphorylation site having the sequence SHIE is found from amino acids 230 to 233. A fourth predicted casein kinase II phosphorylation site having the sequence TDGE is found from amino acids 270 to 273. A fifth predicted casein kinase II phosphorylation site having the sequence SEKD is
30 found from amino acids 287 to 290. A sixth predicted casein kinase II phosphorylation site having the sequence SDPD is found from amino acids 322 to 325. A seventh predicted casein kinase II phosphorylation site having the sequence TSVD is found from amino acids 485 to 488. An eighth predicted casein kinase II phosphorylation site having the sequence TLKD is found from amino acids 530 to 533. A ninth predicted casein kinase II phosphorylation site having the sequence SVQD is
35 found from amino acids 548 to 551. A tenth predicted casein kinase II phosphorylation site having

the sequence SYND is found from amino acids 556 to 559. An eleventh predicted casein kinase II phosphorylation site having the sequence TASE is found from amino acids 593 to 596. A twelfth predicted casein kinase II phosphorylation site having the sequence TMDE is found from amino acids 696 to 669. A thirteenth predicted casein kinase II phosphorylation site having the sequence SGQE is found from amino acids 724 to 727. A fourteenth predicted casein kinase II phosphorylation site having the sequence SLED is found from amino acids 753 to 756. A fifteenth predicted casein kinase II phosphorylation site having the sequence TAME is found from amino acids 801 to 804. A sixteenth predicted casein kinase II phosphorylation site having the sequence SQSE is found from amino acids 859 to 862. A seventeenth predicted casein kinase II phosphorylation site having the sequence STAD is found from amino acids 938 to 941. An eighteenth predicted casein kinase II phosphorylation site having the sequence SHFE is found from amino acids 966 to 969. A nineteenth predicted casein kinase II phosphorylation site having the sequence SSLE is found from amino acids 974 to 977. A twentieth predicted casein kinase II phosphorylation site having the sequence TPTE is found from amino acids 1046 to 1049. A twenty-first predicted casein kinase II phosphorylation site having the sequence SNSD is found from amino acids 1061 to 1064. A twenty-second predicted casein kinase II phosphorylation site having the sequence SKQE is found from amino acids 1133 to 1136.

A predicted tyrosine kinase phosphorylation site having the sequence RPAQDCSSY is found from amino acids 812 to 820.

An predicted N-myristoylation site having the sequence GLLVAW is found from amino acids 6 to 11 of SEQ ID NO:21. A second predicted N-myristoylation site having the sequence GLSLAT is found from amino acids 106 to 111. A third predicted N-myristoylation site having the sequence GGTETR is found from amino acids 236 to 241. A fourth predicted N-myristoylation site having the sequence GIEFAR is found from amino acids 245 to 250. A fifth predicted N-myristoylation site having the sequence GGRKGA is found from amino acids 257 to 262. A sixth predicted N-myristoylation site having the sequence GTNKNE is found from amino acids 354 to 359. A seventh predicted N-myristoylation site having the sequence GLEMSQ is found from amino acids 363 to 368. A eighth predicted N-myristoylation site having the sequence GILLGA is found from amino acids 379 to 384. A ninth predicted N-myristoylation site having the sequence GSYFGS is found from amino acids 477 to 482. A tenth predicted N-myristoylation site having the sequence GSCIAS is found from amino acids 543 to 548. An eleventh predicted N-myristoylation site having the sequence GALGNA is found from amino acids 625 to 630. A twelfth predicted N-myristoylation site having the sequence GIRYNA is found from amino acids 690 to 695. A thirteenth predicted N-myristoylation site having the sequence GQEHCC is found from amino acids 725 to 730. A fourteenth predicted N-myristoylation site having the sequence GNTSCN is found from amino acids

1030 to 1035. A fifteenth predicted N-myristoylation site having the sequence GSTLGG is found from amino acids 1147 to 1152.

A predicted amidation site having the sequence SGKK is found from amino acids 51 to 54 of SEQ ID NO:21. A second predicted amidation site having the sequence GGRK is found from amino acids 257 to 260.

Clone Mouse I2, which encodes murine A259, was deposited with the American Type Culture Collection (10801 University Boulevard, Manassas, VA 20110-2209) on April 2, 1999 and assigned Accession Number 207191. This deposit will be maintained under the terms of the Budapest Treaty on the International Recognition of the Deposit of Microorganisms for the Purposes of Patent Procedure. This deposit was made merely as a convenience for those of skill in the art and is not an admission that a deposit is required under 35 U.S.C. §112.

Figure 6 depicts a hydropathy plot of murine A259. Relatively hydrophobic regions of the protein are shown above the horizontal line, and relatively hydrophilic regions of the protein are below the horizontal line. The cysteine residues (cys) and predicted N-glycosylation sites are indicated by short vertical lines just below the hydropathy trace. The dashed vertical line separates the signal sequence (amino acids 1 to 22 of SEQ ID NO:21; SEQ ID NO:23) on the left from the mature protein (amino acids 23 to 1188 of SEQ ID NO:21; SEQ ID NO:22) on the right. The A259 transmembrane domain is indicated by the section of the plot under which the number 6.4 can be seen, which represents a score assigned to the predicted transmembrane domain. The extracellular domain (SEQ ID NO:24) and cytoplasmic domain (SEQ ID NO:34) are similarly indicated by gray horizontal bars, labeled as A_{out} and A_{in}, respectively.

In situ data of A259 expression in mouse is summarized as follows: During embryogenesis, E13.5 through postnatal day 1.5 tested, all developing bone structures have strong expression. The signal is located mostly at the edge of the developing bones. Beginning at E14.5 the bone marrow space can be seen and is negative for expression. However, some large bones have signal in the bone, not just at the edge. Signal is also observed in the small intestine, diaphragm and to a lesser extent the muscle layer under the skin. The diaphragm and intestine expression pattern is suggestive of smooth muscle. The expression in these tissues is strongest at E15.5 and E16.5 and decreases to almost background levels by P1.5. Adult expression is observed in a small number of tissues examined, including brain, submandibular gland, bladder, colon, small intestine, liver, spleen, and placenta. The signal in positive tissues is very weak in all but the mucosal epithelium of the bladder which has the strongest expression. There is no expression observed in the following tested tissues: spinal cord, eye and Harderian gland, white fat, brown fat, stomach, heart, kidney, adrenal gland, thymus, lymph node, lung, pancreas, skeletal muscle, and testes.

Uses of A259 Nucleic acids, Polypeptides, and Modulators Thereof

As human A259 was originally found in a LPS stimulated human primary osteoblast library, A259 nucleic acids, proteins, and modulators thereof can be used to modulate the proliferation, differentiation, and/or function of cells that form bone matrix, e.g., osteoblasts and osteoclasts, and can be used to modulate the formation of bone matrix. Thus A259 nucleic acids, proteins, and modulators thereof can be used to treat cartilage and bone associated diseases and disorders, and can play a role in bone growth, formation, and remodeling. Examples of cartilage and bone associated diseases and disorders include e.g., bone cancer, achondroplasia, myeloma, fibrous dysplasia, scoliosis, osteoarthritis, osteosarcoma, and osteoporosis.

As murine A259 was originally found in a bone marrow library, A259 nucleic acids, proteins, and modulators thereof can be used to modulate the proliferation, differentiation, and/or function of cells that appear in the bone marrow, e.g., stem cells (e.g., hematopoietic stem cells), and blood cells, e.g., erythrocytes, platelets, and leukocytes. Thus A259 nucleic acids, proteins, and modulators thereof can be used to treat bone marrow, blood, and hematopoietic associated diseases and disorders, e.g., acute myeloid leukemia, hemophilia, leukemia, anemia (e.g., sickle cell anemia), and thalassemia.

As integrin family members play a role in immune response, A259 nucleic acids, proteins, and modulators thereof can be used to treat immune related disorders, e.g., immunodeficiency disorders (e.g., HIV), viral disorders (e.g., infection by HSV), cell growth disorders, e.g., cancers (e.g., carcinoma, lymphoma, e.g., follicular lymphoma), autoimmune disorders (e.g., arthritis, graft rejection (e.g., allograft rejection), T cell autoimmune disorders (e.g., AIDS)), and inflammatory disorders (e.g., bacterial or viral infection, psoriasis, septicemia, cerebral malaria, inflammatory bowel disease, arthritis (e.g., rheumatoid arthritis, osteoarthritis), allergic inflammatory disorders (e.g., asthma, psoriasis)).

As integrin family members play a role in cell growth, survival, proliferation, and migration, A259 nucleic acids, proteins, and modulators thereof can be used to treat apoptotic disorders (e.g., rheumatoid arthritis, systemic lupus erythematosus, insulin-dependent diabetes mellitus) proliferative disorders (e.g., cancers, e.g., B cell cancers stimulated by TNF), and disorders abnormal vascularization (e.g., cancer). In addition, A259 nucleic acids, proteins, and modulators thereof can also be used to promote vascularization (angiogenesis).

As integrins are cell adhesion molecules, A259 nucleic acids, proteins, and modulators thereof can be used to modulate disorders associated with adhesion and migration of cells, e.g., platelet aggregation disorders (e.g., Glanzmann=s thromboasthenia, which is a bleeding disorders characterized by failure of platelet aggregation in response to cell stimuli), inflammatory disorders (e.g., leukocyte adhesion deficiency, which is a disorder associated with impaired migration of

neutrophils to sites of extravascular inflammation), disorders associated with abnormal tissue migration during embryo development, and tumor metastasis.

A259 polypeptides, nucleic acids, and modulators thereof, can also be used to modulate the function, morphology, proliferation and/or differentiation of cells in the tissues in which it is

5 expressed. Such molecules can be used to treat disorders associated with abnormal or aberrant metabolism or function of cells in the tissues in which it is expressed. Tissues in which A259 is expressed, and disorders of which A259 polypeptides, nucleic acids, and modulators thereof can be used to treat, include: bone (see disorders described herein); intestine (e.g., ischemic bowel disease, infective enterocolitis, Crohn=s disease); brain (e.g., cerebral edema, hydrocephalus, brain
10 herniations, iatrogenic disease (due to, e.g., infection, toxins, or drugs)); bladder (e.g., cystitis (bladder infection), incontinence); liver (e.g., jaundice, hepatic failure, hepatic circulatory disorders (e.g., hepatic vein thrombosis and portal vein obstruction and thrombosis), hepatitis (e.g., chronic active hepatitis, acute viral hepatitis, and toxic and drug-induced hepatitis), cirrhosis (e.g., alcoholic cirrhosis, biliary cirrhosis, and hemochromatosis)); spleen (e.g., amyloidosis, Niemann-Pick disease,
15 splenomegaly); placenta (e.g., toxemia of pregnancy (e.g., preeclampsia and eclampsia, placentitis, spontaneous abortion); and neuronal tissue (e.g., epilepsy, muscular dystrophy, and neurodegenerative diseases such as Alzheimer=s disease, Parkinson=s disease, and Huntington=s disease).

A259 is expressed at a higher level in the liver of patients suffering from liver fibrosis than in
20 patients not suffering from liver fibrosis. Liver fibrosis, is caused by chronic injury to the liver arising from, e.g, alcohol abuse, drugs, viral infections (e.g., hepatitis B or hepatitis C), metabolic disorders (excessive iron or copper), autoimmune attack on hepatocytes or the bile duct, and congenital disorders. Liver fibrosis is a reversible condition and injury can be present for months or years before significant scar tissue accumulates. However, liver fibrosis leads to cirrhosis, which is
25 generally not reversible.

The liver has four major components: epithelial cells (hepatocytes), endothelial cells; tissue macrophages (Kupffer cells), and a stellate cells (a type of perivascular mesenchymal cell). In normal liver the space between the epithelium and sinusoidal endothelium is filled with a type of extracellular matrix (ECM) that is somewhat similar to a basement membrane. In fibrosis the ECM
30 undergoes a number of changes. The total content of collagens and other components increases several fold. In addition, there is an increase in fibril-forming collagens. Stellate cells are the primary fibrogenic cell of the liver. In response to injury, quiescent stellate cells are activated and become proliferative, fibrogenic, contractile myofibroblasts. Among the key mediators of these changes are: TGF- β 1, MCP-1, PDGF, ET-1, and MMP-2. Kupffer cells, endothelial cells and
35 hepatocytes produce factors which stimulate stellate cell activation. As noted above, human A259 is

expressed at a high level in activated stellate cells. This result and the increased expression of A259 in clinical liver fibrosis samples, suggest that A259 plays a role in liver fibrosis. Thus, A259 nucleic acids, polypeptides, and modulators thereof can be used to diagnose and treat liver fibrosis as well as other types of fibrosis, e.g., kidney fibrosis or lung fibrosis. Thus, liver fibrosis can be treated using
5 antibodies directed against A259, particularly the extracellular domain of A259, the I domain of A259, or a repeat domain of A259. Also useful are polypeptides which include one or more repeat domains of A259 and/or the I domain of A259.

Tables 1, 2 and 3 below provide summaries of human A259 and murine A259 sequence information.

TABLE 1: Summary of Sequence Information of Human A259 and Murine A259.

Gene	cDNA	Open Reading Frame	Polypeptide	Figure	ATCC Accession No.
Human A259	SEQ ID NO:1	SEQ ID NO:2	SEQ ID NO:3	Figs. 1A-1D	207190
Murine A259	SEQ ID NO:19	SEQ ID NO:20	SEQ ID NO:21	Figs. 4A-4D	207191

TABLE 2: Summary of Domains of Human A259 and Murine A259.

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Protein	Signal Sequence	Mature Protein	Extracellular Domain	Transmembrane Domain	Cytoplasmic Domain
Human A259	aa 1-22 SEQ ID NO:5	aa 23-1188 SEQ ID NO:4	aa 1-1141 SEQ ID NO:6	aa 1142-1164 SEQ ID NO:15	aa 1165-1188 SEQ ID NO:16
Murine A259	aa 1-22 SEQ ID NO:23	aa 23-1188 SEQ ID NO:22	aa 1-1141 SEQ ID NO:24	aa 1142-1164 SEQ ID NO:33	aa 1165-1188 SEQ ID NO:34

TABLE 3: Summary of Domains of Human A259 and Murine A259.

Protein	I Domain	Repeat Domain
Human A259	aa 164-345 SEQ ID NO:7	aa 39-74; 115-157; 367-392; 421-455; 478-516; 540-575; 602-640 SEQ ID NO:8-14
Murine A259	aa 164-345 SEQ ID NO:25	aa 39-74; 115-157; 367-392; 421-455; 478-516; 540-575; 602-640 SEQ ID NO:26-32

10 Various aspects of the invention are described in further detail in the following subsections.

I. Isolated Nucleic Acid Molecules

One aspect of the invention pertains to isolated nucleic acid molecules that encode a polypeptide of the invention or a biologically active portion thereof, as well as nucleic acid molecules sufficient for use as hybridization probes to identify nucleic acid molecules encoding a

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polypeptide of the invention and fragments of such nucleic acid molecules suitable for use as PCR primers for the amplification or mutation of nucleic acid molecules. As used herein, the term "nucleic acid molecule" is intended to include DNA molecules (e.g., cDNA or genomic DNA) and RNA molecules (e.g., mRNA) and analogs of the DNA or RNA generated using nucleotide analogs.

5 The nucleic acid molecule can be single-stranded or double-stranded, but preferably is double-stranded DNA.

An "isolated" nucleic acid molecule is one which is separated from other nucleic acid molecules which are present in the natural source of the nucleic acid molecule. Preferably, an "isolated" nucleic acid molecule is free of sequences (preferably protein encoding sequences) which naturally flank the nucleic acid (i.e., sequences located at the 5' and 3' ends of the nucleic acid) in the genomic DNA of the organism from which the nucleic acid is derived. For example, in various embodiments, the isolated nucleic acid molecule can contain less than about 5 kB, 4 kB, 3 kB, 2 kB, 1 kB, 0.5 kB or 0.1 kB of nucleotide sequences which naturally flank the nucleic acid molecule in genomic DNA of the cell from which the nucleic acid is derived. Moreover, an "isolated" nucleic acid molecule, such as a cDNA molecule, can be substantially free of other cellular material, or culture medium when produced by recombinant techniques, or substantially free of chemical precursors or other chemicals when chemically synthesized. As used herein, the term isolated when referring to a nucleic acid molecule does not include an isolated chromosome.

A nucleic acid molecule of the present invention, e.g., a nucleic acid molecule having the nucleotide sequence of SEQ ID NO:1, 2, 19, or 20, or a complement thereof, can be isolated using standard molecular biology techniques and the sequence information provided herein. Using all or a portion of the nucleic acid sequences of SEQ ID NO:1, 2, 19, or 20 as a hybridization probe, nucleic acid molecules of the invention can be isolated using standard hybridization and cloning techniques (e.g., as described in Sambrook et al., eds., *Molecular Cloning: A Laboratory Manual*, 2nd ed., Cold Spring Harbor Laboratory, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, 1989).

A nucleic acid molecule of the invention can be amplified using cDNA, mRNA or genomic DNA as a template and appropriate oligonucleotide primers according to standard PCR amplification techniques. The nucleic acid so amplified can be cloned into an appropriate vector and characterized by DNA sequence analysis. Furthermore, oligonucleotides corresponding to all or a portion of a nucleic acid molecule of the invention can be prepared by standard synthetic techniques, e.g., using an automated DNA synthesizer.

In another preferred embodiment, an isolated nucleic acid molecule of the invention comprises a nucleic acid molecule which is a complement of the nucleotide sequence of SEQ ID NO:1, 2, 19, or 20, or a portion thereof. A nucleic acid molecule which is complementary to a given

nucleotide sequence is one which is sufficiently complementary to the given nucleotide sequence that it can hybridize to the given nucleotide sequence thereby forming a stable duplex.

Moreover, a nucleic acid molecule of the invention can comprise only a portion of a nucleic acid sequence encoding a full length polypeptide of the invention for example, a fragment which can be used as a probe or primer or a fragment encoding a biologically active portion of a polypeptide of the invention. The nucleotide sequence determined from the cloning one gene allows for the generation of probes and primers designed for use in identifying and/or cloning homologues in other cell types, e.g., from other tissues, as well as homologues from other mammals. The probe/primer typically comprises substantially purified oligonucleotide. The oligonucleotide typically comprises a region of nucleotide sequence that hybridizes under stringent conditions to at least about 12, preferably about 25, more preferably about 50, 75, 100, 125, 150, 175, 200, 250, 300, 350 or 400 consecutive nucleotides of the sense or anti-sense sequence of SEQ ID NO:1, 2, 19, or 20 or of a naturally occurring mutant of SEQ ID NO:1, 2, 19, or 20.

Probes based on the sequence of a nucleic acid molecule of the invention can be used to detect transcripts or genomic sequences encoding the same protein molecule encoded by a selected nucleic acid molecule. The probe comprises a label group attached thereto, e.g., a radioisotope, a fluorescent compound, an enzyme, or an enzyme co-factor. Such probes can be used as part of a diagnostic test kit for identifying cells or tissues which mis-express the protein, such as by measuring levels of a nucleic acid molecule encoding the protein in a sample of cells from a subject, e.g., detecting mRNA levels or determining whether a gene encoding the protein has been mutated or deleted.

A nucleic acid fragment encoding a biologically active portion of a polypeptide of the invention can be prepared by isolating a portion of any of SEQ ID NO:2 or 20, expressing the encoded portion of the polypeptide protein (e.g., by recombinant expression *in vitro*) and assessing the activity of the encoded portion of the polypeptide.

The invention further encompasses nucleic acid molecules that differ from the nucleotide sequence of SEQ ID NO:1, 2, 19, or 20, due to degeneracy of the genetic code and thus encode the same protein as that encoded by the nucleotide sequence of SEQ ID NO:2 or 20.

In addition to the nucleotide sequences of SEQ ID NO:2 and 20, it will be appreciated by those skilled in the art that DNA sequence polymorphisms that lead to changes in the amino acid sequence may exist within a population (e.g., the human population). Such genetic polymorphisms may exist among individuals within a population due to natural allelic variation. An allele is one of a group of genes which occur alternatively at a given genetic locus. As used herein, the phrase allelic variant refers to a nucleotide sequence which occurs at a given locus or to a polypeptide encoded by the nucleotide sequence. As used herein, the terms "gene" and "recombinant gene" refer to nucleic

acid molecules comprising an open reading frame encoding a polypeptide of the invention. Such natural allelic variations can typically result in 1-5% variance in the nucleotide sequence of a given gene. Alternative alleles can be identified by sequencing the gene of interest in a number of different individuals. This can be readily carried out by using hybridization probes to identify the same genetic locus in a variety of individuals. Any and all such nucleotide variations and resulting amino acid polymorphisms or variations that are the result of natural allelic variation and that do not alter the functional activity are intended to be within the scope of the invention.

Moreover, nucleic acid molecules encoding proteins of the invention from other species (homologues), which have a nucleotide sequence which differs from that of the human protein described herein are intended to be within the scope of the invention. Nucleic acid molecules corresponding to natural allelic variants and homologues of a cDNA of the invention can be isolated based on their identity to the human nucleic acid molecule disclosed herein using the human cDNAs, or a portion thereof, as a hybridization probe according to standard hybridization techniques under stringent hybridization conditions. For example, a cDNA encoding a soluble form of a membrane-bound protein of the invention isolated based on its hybridization to a nucleic acid molecule encoding all or part of the membrane-bound form. Likewise, a cDNA encoding a membrane-bound form can be isolated based on its hybridization to a nucleic acid molecule encoding all or part of the soluble form.

Accordingly, in another embodiment, an isolated nucleic acid molecule of the invention is at least 400 (450, 500, 550, 600, 650, 700, 800, 900, 1000, 2000, or 3000) nucleotides in length and hybridizes under stringent conditions to the nucleic acid molecule comprising the nucleotide sequence, preferably the coding sequence, of SEQ ID NO:1 or 19, or a complement thereof.

As used herein, the term "hybridizes under stringent conditions" is intended to describe conditions for hybridization and washing under which nucleotide sequences at least 60% (65%, 70%, preferably 75%) identical to each other typically remain hybridized to each other. Such stringent conditions are known to those skilled in the art and can be found in *Current Protocols in Molecular Biology*, John Wiley & Sons, N.Y. (1989), 6.3.1-6.3.6. A preferred, non-limiting example of stringent hybridization conditions are hybridization in 6X sodium chloride/sodium citrate (SSC) at about 45EC, followed by one or more washes in 0.2 X SSC, 0.1% SDS at 50-65EC. Preferably, an isolated nucleic acid molecule of the invention that hybridizes under stringent conditions to the sequence of SEQ ID NO:1, 2, 19, or 20, or a complement thereof, corresponds to a naturally-occurring nucleic acid molecule. As used herein, a "naturally-occurring" nucleic acid molecule refers to an RNA or DNA molecule having a nucleotide sequence that occurs in nature (e.g., encodes a natural protein).

In addition to naturally-occurring allelic variants of a nucleic acid molecule of the invention sequence that may exist in the population, the skilled artisan will further appreciate that changes can be introduced by mutation thereby leading to changes in the amino acid sequence of the encoded protein, without altering the biological activity of the protein. For example, one can make nucleotide
5 substitutions leading to amino acid substitutions at "non-essential" amino acid residues. A "non-essential" amino acid residue is a residue that can be altered from the wild-type sequence without altering the biological activity, whereas an "essential" amino acid residue is required for biological activity. For example, amino acid residues that are not conserved or only semi-conserved among homologues of various species may be non-essential for activity and thus would be likely targets for
10 alteration. Alternatively, amino acid residues that are conserved among the homologues of various species (e.g., murine and human) may be essential for activity and thus would not be likely targets for alteration.

Accordingly, another aspect of the invention pertains to nucleic acid molecules encoding a polypeptide of the invention that contain changes in amino acid residues that are not essential for
15 activity. Such polypeptides differ in amino acid sequence from SEQ ID NO:3 or 21, yet retain biological activity. In one embodiment, the isolated nucleic acid molecule includes a nucleotide sequence encoding a protein that includes an amino acid sequence that is at least about 44% (or 45%, 50%, 55%, 60%, 65%, 75%, 85%, 95%, or 98%) identical to the amino acid sequence of SEQ ID NO:3 or 21.

An isolated nucleic acid molecule encoding a variant protein can be created by introducing one or more nucleotide substitutions, additions or deletions into the nucleotide sequence of SEQ ID NO:1, 2, 19, or 20 such that one or more amino acid substitutions, additions or deletions are introduced into the encoded protein. Mutations can be introduced by standard techniques, such as site-directed mutagenesis and PCR-mediated mutagenesis. Preferably, conservative amino acid
20 substitutions are made at one or more predicted non-essential amino acid residues. A "conservative amino acid substitution" is one in which the amino acid residue is replaced with an amino acid residue having a similar side chain. Families of amino acid residues having similar side chains have been defined in the art. These families include amino acids with basic side chains (e.g., lysine, arginine, histidine), acidic side chains (e.g., aspartic acid, glutamic acid), uncharged polar side chains
25 (e.g., glycine, asparagine, glutamine, serine, threonine, tyrosine, cysteine), nonpolar side chains (e.g., alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine, tryptophan), beta-branched side chains (e.g., threonine, valine, isoleucine) and aromatic side chains (e.g., tyrosine, phenylalanine, tryptophan, histidine). Alternatively, mutations can be introduced randomly along all or part of the coding sequence, such as by saturation mutagenesis, and the resultant mutants can be
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screened for biological activity to identify mutants that retain activity. Following mutagenesis, the encoded protein can be expressed recombinantly and the activity of the protein can be determined.

In a preferred embodiment, a mutant polypeptide that is a variant of a polypeptide of the invention can be assayed for: (1) the ability to form protein:protein interactions with proteins in a signaling pathway of the polypeptide of the invention; (2) the ability to bind a ligand of the polypeptide of the invention; or (3) the ability to bind to an intracellular target protein of the polypeptide of the invention. In yet another preferred embodiment, the mutant polypeptide can be assayed for the ability to modulate cellular proliferation, cellular migration or chemotaxis, or cellular differentiation.

The present invention encompasses antisense nucleic acid molecules, i.e., molecules which are complementary to a sense nucleic acid encoding a polypeptide of the invention, e.g., complementary to the coding strand of a double-stranded cDNA molecule or complementary to an mRNA sequence. Accordingly, an antisense nucleic acid can hydrogen bond to a sense nucleic acid. The antisense nucleic acid can be complementary to an entire coding strand, or to only a portion thereof, e.g., all or part of the protein coding region (or open reading frame). An antisense nucleic acid molecule can be antisense to all or part of a non-coding region of the coding strand of a nucleotide sequence encoding a polypeptide of the invention. The non-coding regions ("5' and 3' untranslated regions") are the 5' and 3' sequences which flank the coding region and are not translated into amino acids.

An antisense oligonucleotide can be, for example, about 5, 10, 15, 20, 25, 30, 35, 40, 45 or 50 nucleotides or more in length. An antisense nucleic acid of the invention can be constructed using chemical synthesis and enzymatic ligation reactions using procedures known in the art. For example, an antisense nucleic acid (e.g., an antisense oligonucleotide) can be chemically synthesized using naturally occurring nucleotides or variously modified nucleotides designed to increase the biological stability of the molecules or to increase the physical stability of the duplex formed between the antisense and sense nucleic acids, e.g., phosphorothioate derivatives and acridine substituted nucleotides can be used. Examples of modified nucleotides which can be used to generate the antisense nucleic acid include 5-fluorouracil, 5-bromouracil, 5-chlorouracil, 5-iodouracil, hypoxanthine, xanthine, 4-acetylcytosine, 5-(carboxyhydroxymethyl) uracil, 5-carboxymethylaminomethyl-2-thiouridine, 5-carboxymethylaminomethyluracil, dihydrouracil, beta-D-galactosylqueosine, inosine, N6-isopentenyladenine, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine, 2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-adenine, 7-methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5'-methoxycarboxymethyluracil, 5-methoxyuracil, 2-methylthio-N6-isopentenyladenine, uracil-5-oxyacetic acid (v), wybutoxosine, pseudouracil, queosine, 2-

thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil, uracil-5-oxyacetic acid methylester, uracil-5-oxyacetic acid (v), 5-methyl-2-thiouracil, 3-(3-amino-3-predicted N-2-carboxypropyl) uracil, (acp3)w, and 2,6-diaminopurine. Alternatively, the antisense nucleic acid can be produced biologically using an expression vector into which a nucleic acid has been subcloned in an antisense orientation (i.e., RNA transcribed from the inserted nucleic acid will be of an antisense orientation to a target nucleic acid of interest, described further in the following subsection).

The antisense nucleic acid molecules of the invention are typically administered to a subject or generated *in situ* such that they hybridize with or bind to cellular mRNA and/or genomic DNA encoding a selected polypeptide of the invention to thereby inhibit expression, e.g., by inhibiting transcription and/or translation. The hybridization can be by conventional nucleotide complementarity to form a stable duplex, or, for example, in the case of an antisense nucleic acid molecule which binds to DNA duplexes, through specific interactions in the major groove of the double helix. An example of a route of administration of antisense nucleic acid molecules of the invention includes direct injection at a tissue site. Alternatively, antisense nucleic acid molecules can be modified to target selected cells and then administered systemically. For example, for systemic administration, antisense molecules can be modified such that they specifically bind to receptors or antigens expressed on a selected cell surface, e.g., by linking the antisense nucleic acid molecules to peptides or antibodies which bind to cell surface receptors or antigens. The antisense nucleic acid molecules can also be delivered to cells using the vectors described herein. To achieve sufficient intracellular concentrations of the antisense molecules, vector constructs in which the antisense nucleic acid molecule is placed under the control of a strong pol II or pol III promoter are preferred.

An antisense nucleic acid molecule of the invention can be an α -anomeric nucleic acid molecule. An α -anomeric nucleic acid molecule forms specific double-stranded hybrids with complementary RNA in which, contrary to the usual β -units, the strands run parallel to each other (Gaultier et al. (1987) *Nucleic Acids Res.* 15:6625-6641). The antisense nucleic acid molecule can also comprise a 2'-o-methylribonucleotide (Inoue et al. (1987) *Nucleic Acids Res.* 15:6131-6148) or a chimeric RNA-DNA analogue (Inoue et al. (1987) *FEBS Lett.* 215:327-330).

The invention also encompasses ribozymes. Ribozymes are catalytic RNA molecules with ribonuclease activity which are capable of cleaving a single-stranded nucleic acid, such as an mRNA, to which they have a complementary region. Thus, ribozymes (e.g., hammerhead ribozymes (described in Haselhoff and Gerlach (1988) *Nature* 334:585-591)) can be used to catalytically cleave mRNA transcripts to thereby inhibit translation of the protein encoded by the mRNA. A ribozyme having specificity for a nucleic acid molecule encoding a polypeptide of the invention can be designed based upon the nucleotide sequence of a cDNA disclosed herein. For example, a derivative of a *Tetrahymena* L-19 IVS RNA can be constructed in which the nucleotide sequence of the active

site is complementary to the nucleotide sequence to be cleaved in a Cech et al. U.S. Patent No. 4,987,071; and Cech et al. U.S. Patent No. 5,116,742. Alternatively, an mRNA encoding a polypeptide of the invention can be used to select a catalytic RNA having a specific ribonuclease activity from a pool of RNA molecules. *See, e.g.,* Bartel and Szostak (1993) *Science* 261:1411-1418.

5 The invention also encompasses nucleic acid molecules which form triple helical structures. For example, expression of a polypeptide of the invention can be inhibited by targeting nucleotide sequences complementary to the regulatory region of the gene encoding the polypeptide (e.g., the promoter and/or enhancer) to form triple helical structures that prevent transcription of the gene in target cells. *See generally* Helene (1991) *Anticancer Drug Des.* 6(6):569-84; Helene (1992) *Ann.*
10 *N.Y. Acad. Sci.* 660:27-36; and Maher (1992) *Bioassays* 14(12):807-15.

In various embodiments, the nucleic acid molecules of the invention can be modified at the base moiety, sugar moiety or phosphate backbone to improve, e.g., the stability, hybridization, or solubility of the molecule. For example, the deoxyribose phosphate backbone of the nucleic acids can be modified to generate peptide nucleic acids (*see* Hyrup et al. (1996) *Bioorganic & Medicinal*
15 *Chemistry* 4(1): 5-23). As used herein, the terms "peptide nucleic acids" or "PNAs" refer to nucleic acid mimics, e.g., DNA mimics, in which the deoxyribose phosphate backbone is replaced by a pseudopeptide backbone and only the four natural nucleobases are retained. The neutral backbone of PNAs has been shown to allow for specific hybridization to DNA and RNA under conditions of low ionic strength. The synthesis of PNA oligomers can be performed using standard solid phase peptide
20 synthesis protocols as described in Hyrup et al. (1996), *supra*; Perry-O'Keefe et al. (1996) *Proc. Natl. Acad. Sci. USA* 93: 14670-675.

PNAs can be used in therapeutic and diagnostic applications. For example, PNAs can be used as antisense or antigene agents for sequence-specific modulation of gene expression by, e.g., inducing transcription or translation arrest or inhibiting replication. PNAs can also be used, e.g., in
25 the analysis of single base pair mutations in a gene by, e.g., PNA directed PCR clamping; as artificial restriction enzymes when used in combination with other enzymes, e.g., S1 nucleases (Hyrup (1996), *supra*; or as probes or primers for DNA sequence and hybridization (Hyrup (1996), *supra*; Perry-O'Keefe et al. (1996) *Proc. Natl. Acad. Sci. USA* 93: 14670-675).

In another embodiment, PNAs can be modified, e.g., to enhance their stability or cellular
30 uptake, by attaching lipophilic or other helper groups to PNA, by the formation of PNA-DNA chimeras, or by the use of liposomes or other techniques of drug delivery known in the art. For example, PNA-DNA chimeras can be generated which may combine the advantageous properties of PNA and DNA. Such chimeras allow DNA recognition enzymes, e.g., RNase H and DNA polymerases, to interact with the DNA portion while the PNA portion would provide high binding
35 affinity and specificity. PNA-DNA chimeras can be linked using linkers of appropriate lengths

selected in terms of base stacking, number of bonds between the nucleobases, and orientation (Hyrup (1996), *supra*). The synthesis of PNA-DNA chimeras can be performed as described in Hyrup (1996), *supra*, and Finn et al. (1996) *Nucleic Acids Res.* 24(17):3357-63. For example, a DNA chain can be synthesized on a solid support using standard phosphoramidite coupling chemistry and modified nucleoside analogs. Compounds such as 5'-(4-methoxytrityl)amino-5'-deoxy-thymidine phosphoramidite can be used as a link between the PNA and the 5' end of DNA (Mag et al. (1989) *Nucleic Acids Res.* 17:5973-88). PNA monomers are then coupled in a stepwise manner to produce a chimeric molecule with a 5' PNA segment and a 3' DNA segment (Finn et al. (1996) *Nucleic Acids Res.* 24(17):3357-63). Alternatively, chimeric molecules can be synthesized with a 5' DNA segment and a 3' PNA segment (Peterser et al. (1975) *Bioorganic Med. Chem. Lett.* 5:1119-1124).

In other embodiments, the oligonucleotide may include other appended groups such as peptides (e.g., for targeting host cell receptors *in vivo*), or agents facilitating transport across the cell membrane (see, e.g., Letsinger et al. (1989) *Proc. Natl. Acad. Sci. USA* 86:6553-6556; Lemaitre et al. (1987) *Proc. Natl. Acad. Sci. USA* 84:648-652; PCT Publication No. W0 88/09810) or the blood-brain barrier (see, e.g., PCT Publication No. W0 89/10134). In addition, oligonucleotides can be modified with hybridization-triggered cleavage agents (see, e.g., Krol et al. (1988) *Bio/Techniques* 6:958-976) or intercalating agents (see, e.g., Zon (1988) *Pharm. Res.* 5:539-549). To this end, the oligonucleotide may be conjugated to another molecule, e.g., a peptide, hybridization triggered cross-linking agent, transport agent, hybridization-triggered cleavage agent, etc.

II. Isolated Proteins and Antibodies

One aspect of the invention pertains to isolated proteins, and biologically active portions thereof, as well as polypeptide fragments suitable for use as immunogens to raise antibodies directed against a polypeptide of the invention. In one embodiment, the native polypeptide can be isolated from cells or tissue sources by an appropriate purification scheme using standard protein purification techniques. In another embodiment, polypeptides of the invention are produced by recombinant DNA techniques. Alternative to recombinant expression, a polypeptide of the invention can be synthesized chemically using standard peptide synthesis techniques.

An "isolated" or "purified" protein or biologically active portion thereof is substantially free of cellular material or other contaminating proteins from the cell or tissue source from which the protein is derived, or substantially free of chemical precursors or other chemicals when chemically synthesized. The language "substantially free of cellular material" includes preparations of protein in which the protein is separated from cellular components of the cells from which it is isolated or recombinantly produced. Thus, protein that is substantially free of cellular material includes preparations of protein having less than about 30%, 20%, 10%, or 5% (by dry weight) of

heterologous protein (also referred to herein as a "contaminating protein"). When the protein or biologically active portion thereof is recombinantly produced, it is also preferably substantially free of culture medium, i.e., culture medium represents less than about 20%, 10%, or 5% of the volume of the protein preparation. When the protein is produced by chemical synthesis, it is preferably substantially free of chemical precursors or other chemicals, i.e., it is separated from chemical precursors or other chemicals which are involved in the synthesis of the protein. Accordingly such preparations of the protein have less than about 30%, 20%, 10%, 5% (by dry weight) of chemical precursors or compounds other than the polypeptide of interest.

Biologically active portions of a polypeptide of the invention include polypeptides comprising amino acid sequences sufficiently identical to or derived from the amino acid sequence of the protein (e.g., the amino acid sequence shown in any of SEQ ID NO:3, 4, 21, or 22), which include fewer amino acids than the full length protein, and exhibit at least one activity of the corresponding full-length protein. Typically, biologically active portions comprise a domain or motif with at least one activity of the corresponding protein. A biologically active portion of a protein of the invention can be a polypeptide which is, for example, 10, 25, 50, 100 or more amino acids in length. Moreover, other biologically active portions, in which other regions of the protein are deleted, can be prepared by recombinant techniques and evaluated for one or more of the functional activities of the native form of a polypeptide of the invention.

Preferred polypeptides have the amino acid sequence of SEQ ID NO:3, 4, 21, or 22. Other useful proteins are substantially identical (e.g., at least about 45%, preferably 55%, 65%, 75%, 85%, 95%, or 99%) to any of SEQ ID NO:3, 4, 21, or 22, and retain the functional activity of the protein of the corresponding naturally-occurring protein yet differ in amino acid sequence due to natural allelic variation or mutagenesis.

To determine the percent identity of two amino acid sequences or of two nucleic acids, the sequences are aligned for optimal comparison purposes (e.g., gaps can be introduced in the sequence of a first amino acid or nucleic acid sequence for optimal alignment with a second amino or nucleic acid sequence). The amino acid residues or nucleotides at corresponding amino acid positions or nucleotide positions are then compared. When a position in the first sequence is occupied by the same amino acid residue or nucleotide as the corresponding position in the second sequence, then the molecules are identical at that position. The percent identity between the two sequences is a function of the number of identical positions shared by the sequences (i.e., % identity = # of identical positions/total # of positions (e.g., overlapping positions) x 100). In one embodiment, the two sequences are the same length.

The determination of percent identity between two sequences can be accomplished using a mathematical algorithm. A preferred, non-limiting example of a mathematical algorithm utilized for

the comparison of two sequences is the algorithm of Karlin and Altschul (1990) *Proc. Natl. Acad. Sci. USA* 87:2264-2268, modified as in Karlin and Altschul (1993) *Proc. Natl. Acad. Sci. USA* 90:5873-5877. Such an algorithm is incorporated into the NBLAST and XBLAST programs of Altschul, et al. (1990) *J. Mol. Biol.* 215:403-410. BLAST nucleotide searches can be performed with the NBLAST program, score = 100, wordlength = 12 to obtain nucleotide sequences homologous to a nucleic acid molecules of the invention. BLAST protein searches can be performed with the XBLAST program, score = 50, wordlength = 3 to obtain amino acid sequences homologous to a protein molecules of the invention. To obtain gapped alignments for comparison purposes, Gapped BLAST can be utilized as described in Altschul et al. (1997) *Nucleic Acids Res.* 25:3389-3402. Alternatively, PSI-Blast can be used to perform an iterated search which detects distant relationships between molecules (*Id.*). When utilizing BLAST, Gapped BLAST, and PSI-Blast programs, the default parameters of the respective programs (e.g., XBLAST and NBLAST) can be used. See <http://www.ncbi.nlm.nih.gov>.

Another preferred, non-limiting example of a mathematical algorithm utilized for the comparison of sequences is the algorithm of Myers and Miller, CABIOS (1989). Such an algorithm is incorporated into the ALIGN program (version 2.0) which is part of the CGC sequence alignment software package. When utilizing the ALIGN program for comparing amino acid sequences, a PAM120 weight residue table, a gap length penalty of 12, and a gap penalty of 4 can be used. Additional algorithms for sequence analysis are known in the art and include ADVANCE and ADAM as described in Torellis and Robotti (1994) *Comput. Appl. Biosci.*, 10:3-5; and FASTA described in Pearson and Lipman (1988) *Proc. Natl. Acad. Sci.* 85:2444-8. Within FASTA, ktup is a control option that sets the sensitivity and speed of the search. If ktup=2, similar regions in the two sequences being compared are found by looking at pairs of aligned residues; if ktup=1, single aligned amino acids are examined. ktup can be set to 2 or 1 for protein sequences, or from 1 to 6 for DNA sequences. The default if ktup is not specified is 2 for proteins and 6 for DNA. For a further description of FASTA parameters, see <http://bioweb.pasteur.fr/docs/man/man/fasta.1.html#sect2>, the contents of which are incorporated herein by reference.

The percent identity between two sequences can be determined using techniques similar to those described above, with or without allowing gaps. In calculating percent identity, typically only exact matches are counted.

The invention also provides chimeric or fusion proteins. As used herein, a "chimeric protein" or "fusion protein" comprises all or part (preferably biologically active) of a polypeptide of the invention operably linked to a heterologous polypeptide (i.e., a polypeptide other than the same polypeptide of the invention). Within the fusion protein, the term "operably linked" is intended to indicate that the polypeptide of the invention and the heterologous polypeptide are fused in-frame to

each other. The heterologous polypeptide can be fused to the predicted N-terminus or C-terminus of the polypeptide of the invention.

One useful fusion protein is a GST fusion protein in which the polypeptide of the invention is fused to the C-terminus of GST sequences. Such fusion proteins can facilitate the purification of a recombinant polypeptide of the invention.

In another embodiment, the fusion protein contains a heterologous signal sequence at its predicted N-terminus. For example, the native signal sequence of a polypeptide of the invention can be removed and replaced with a signal sequence from another protein. For example, the gp67 secretory sequence of the baculovirus envelope protein can be used as a heterologous signal sequence (*Current Protocols in Molecular Biology*, Ausubel et al., eds., John Wiley & Sons, 1992). Other examples of eukaryotic heterologous signal sequences include the secretory sequences of melittin and human placental alkaline phosphatase (Stratagene; La Jolla, California). In yet another example, useful prokaryotic heterologous signal sequences include the phoA secretory signal (Sambrook et al., *supra*) and the protein A secretory signal (Pharmacia Biotech; Piscataway, New Jersey).

In yet another embodiment, the fusion protein is an immunoglobulin fusion protein in which all or part of a polypeptide of the invention is fused to sequences derived from a member of the immunoglobulin protein family. The immunoglobulin fusion proteins of the invention can be incorporated into pharmaceutical compositions and administered to a subject to inhibit an interaction between a ligand (soluble or membrane-bound) and a protein on the surface of a cell (receptor), to thereby suppress signal transduction *in vivo*. The immunoglobulin fusion protein can be used to affect the bioavailability of a cognate ligand of a polypeptide of the invention. Inhibition of ligand/receptor interaction may be useful therapeutically, both for treating proliferative and differentiative disorders and for modulating (e.g., promoting or inhibiting) cell survival. Moreover, the immunoglobulin fusion proteins of the invention can be used as immunogens to produce antibodies directed against a polypeptide of the invention in a subject, to purify ligands and in screening assays to identify molecules which inhibit the interaction of receptors with ligands.

Chimeric and fusion proteins of the invention can be produced by standard recombinant DNA techniques. In another embodiment, the fusion gene can be synthesized by conventional techniques including automated DNA synthesizers. Alternatively, PCR amplification of gene fragments can be carried out using anchor primers which give rise to complementary overhangs between two consecutive gene fragments which can subsequently be annealed and reamplified to generate a chimeric gene sequence (*see, e.g.,* Ausubel et al., *supra*). Moreover, many expression vectors are commercially available that already encode a fusion moiety (e.g., a GST polypeptide). A nucleic acid encoding a polypeptide of the invention can be cloned into such an expression vector such that the fusion moiety is linked in-frame to the polypeptide of the invention.

A signal sequence of a polypeptide of the invention (SEQ ID NO:5 and 19) can be used to facilitate secretion and isolation of the secreted protein or other proteins of interest. Signal sequences are typically characterized by a core of hydrophobic amino acids which are generally cleaved from the mature protein during secretion in one or more cleavage events. Such signal peptides contain processing sites that allow cleavage of the signal sequence from the mature proteins as they pass through the secretory pathway. Thus, the invention pertains to the described polypeptides having a signal sequence, as well as to the signal sequence itself and to the polypeptide in the absence of the signal sequence (i.e., the cleavage products). In one embodiment, a nucleic acid sequence encoding a signal sequence of the invention can be operably linked in an expression vector to a protein of interest, such as a protein which is ordinarily not secreted or is otherwise difficult to isolate. The signal sequence directs secretion of the protein, such as from a eukaryotic host into which the expression vector is transformed, and the signal sequence is subsequently or concurrently cleaved. The protein can then be readily purified from the extracellular medium by art recognized methods. Alternatively, the signal sequence can be linked to the protein of interest using a sequence which facilitates purification, such as with a GST domain.

In another embodiment, the signal sequences of the present invention can be used to identify regulatory sequences, e.g., promoters, enhancers, repressors. Since signal sequences are the most amino-terminal sequences of a peptide, it is expected that the nucleic acids which flank the signal sequence on its amino-terminal side will be regulatory sequences which affect transcription. Thus, a nucleotide sequence which encodes all or a portion of a signal sequence can be used as a probe to identify and isolate signal sequences and their flanking regions, and these flanking regions can be studied to identify regulatory elements therein.

The present invention also pertains to variants of the polypeptides of the invention. Such variants have an altered amino acid sequence which can function as either agonists (mimetics) or as antagonists. Variants can be generated by mutagenesis, e.g., discrete point mutation or truncation. An agonist can retain substantially the same, or a subset, of the biological activities of the naturally occurring form of the protein. An antagonist of a protein can inhibit one or more of the activities of the naturally occurring form of the protein by, for example, competitively binding to a downstream or upstream member of a cellular signaling cascade which includes the protein of interest. Thus, specific biological effects can be elicited by treatment with a variant of limited function. Treatment of a subject with a variant having a subset of the biological activities of the naturally occurring form of the protein can have fewer side effects in a subject relative to treatment with the naturally occurring form of the protein.

Variants of a protein of the invention which function as either agonists (mimetics) or as antagonists can be identified by screening combinatorial libraries of mutants, e.g., truncation

mutants, of the protein of the invention for agonist or antagonist activity. In one embodiment, a variegated library of variants is generated by combinatorial mutagenesis at the nucleic acid level and is encoded by a variegated gene library. A variegated library of variants can be produced by, for example, enzymatically ligating a mixture of synthetic oligonucleotides into gene sequences such that a degenerate set of potential protein sequences is expressible as individual polypeptides, or alternatively, as a set of larger fusion proteins (e.g., for phage display). There are a variety of methods which can be used to produce libraries of potential variants of the polypeptides of the invention from a degenerate oligonucleotide sequence. Methods for synthesizing degenerate oligonucleotides are known in the art (*see, e.g.*, Narang (1983) *Tetrahedron* 39:3; Itakura et al. (1984) *Annu. Rev. Biochem.* 53:323; Itakura et al. (1984) *Science* 198:1056; Ike et al. (1983) *Nucleic Acid Res.* 11:477).

In addition, libraries of fragments of the coding sequence of a polypeptide of the invention can be used to generate a variegated population of polypeptides for screening and subsequent selection of variants. For example, a library of coding sequence fragments can be generated by treating a double stranded PCR fragment of the coding sequence of interest with a nuclease under conditions wherein nicking occurs only about once per molecule, denaturing the double stranded DNA, renaturing the DNA to form double stranded DNA which can include sense/antisense pairs from different nicked products, removing single stranded portions from reformed duplexes by treatment with S1 nuclease, and ligating the resulting fragment library into an expression vector. By this method, an expression library can be derived which encodes predicted N-terminal and internal fragments of various sizes of the protein of interest.

Several techniques are known in the art for screening gene products of combinatorial libraries made by point mutations or truncation, and for screening cDNA libraries for gene products having a selected property. The most widely used techniques, which are amenable to high through-put analysis, for screening large gene libraries typically include cloning the gene library into replicable expression vectors, transforming appropriate cells with the resulting library of vectors, and expressing the combinatorial genes under conditions in which detection of a desired activity facilitates isolation of the vector encoding the gene whose product was detected. Recursive ensemble mutagenesis (REM), a technique which enhances the frequency of functional mutants in the libraries, can be used in combination with the screening assays to identify variants of a protein of the invention (Arkin and Yourvan (1992) *Proc. Natl. Acad. Sci. USA* 89:7811-7815; Delgrave et al. (1993) *Protein Engineering* 6(3):327-331).

An isolated polypeptide of the invention, or a fragment thereof, can be used as an immunogen to generate antibodies using standard techniques for polyclonal and monoclonal antibody preparation. The full-length polypeptide or protein can be used or, alternatively, the invention

provides antigenic peptide fragments for use as immunogens. The antigenic peptide of a protein of the invention comprises at least 8 (preferably 10, 15, 20, or 30) amino acid residues of the amino acid sequence of SEQ ID NO:3 or 21, and encompasses an epitope of the protein such that an antibody raised against the peptide forms a specific immune complex with the protein.

5 Preferred epitopes encompassed by the antigenic peptide are regions that are located on the surface of the protein, e.g., hydrophilic regions. Figures 2 and 6 are hydropathy plots of the proteins of the invention. These plots or similar analyses can be used to identify hydrophilic regions.

An immunogen typically is used to prepare antibodies by immunizing a suitable subject, (e.g., rabbit, goat, mouse or other mammal). An appropriate immunogenic preparation can contain, 10 for example, recombinantly expressed or chemically synthesized polypeptide. The preparation can further include an adjuvant, such as Freund's complete or incomplete adjuvant, or similar immunostimulatory agent.

Accordingly, another aspect of the invention pertains to antibodies directed against a polypeptide of the invention. The term "antibody" as used herein refers to immunoglobulin 15 molecules and immunologically active portions of immunoglobulin molecules, i.e., molecules that contain an antigen binding site which specifically binds an antigen, such as a polypeptide of the invention. A molecule which specifically binds to a given polypeptide of the invention is a molecule which binds the polypeptide, but does not substantially bind other molecules in a sample, e.g., a biological sample, which naturally contains the polypeptide. Examples of immunologically active 20 portions of immunoglobulin molecules include F(ab) and F(ab)₂ fragments which can be generated by treating the antibody with an enzyme such as pepsin. The invention provides polyclonal and monoclonal antibodies. The term "monoclonal antibody" or "monoclonal antibody composition", as used herein, refers to a population of antibody molecules that contain only one species of an antigen binding site capable of immunoreacting with a particular epitope.

25 Polyclonal antibodies can be prepared as described above by immunizing a suitable subject with a polypeptide of the invention as an immunogen. The antibody titer in the immunized subject can be monitored over time by standard techniques, such as with an enzyme linked immunosorbent assay (ELISA) using immobilized polypeptide. If desired, the antibody molecules can be isolated from the mammal (e.g., from the blood) and further purified by well-known techniques, such as 30 protein A chromatography to obtain the IgG fraction. At an appropriate time after immunization, e.g., when the specific antibody titers are highest, antibody-producing cells can be obtained from the subject and used to prepare monoclonal antibodies by standard techniques, such as the hybridoma technique originally described by Kohler and Milstein (1975) *Nature* 256:495-497, the human B cell hybridoma technique (Kozbor et al. (1983) *Immunol. Today* 4:72), the EBV-hybridoma technique 35 (Cole et al. (1985), *Monoclonal Antibodies and Cancer Therapy*, Alan R. Liss, Inc., pp. 77-96) or

trionoma techniques. The technology for producing hybridomas is well known (*see generally Current Protocols in Immunology* (1994) Coligan et al. (eds.) John Wiley & Sons, Inc., New York, NY). Hybridoma cells producing a monoclonal antibody of the invention are detected by screening the hybridoma culture supernatants for antibodies that bind the polypeptide of interest, e.g., using a standard ELISA assay.

Alternative to preparing monoclonal antibody-secreting hybridomas, a monoclonal antibody directed against a polypeptide of the invention can be identified and isolated by screening a recombinant combinatorial immunoglobulin library (e.g., an antibody phage display library) with the polypeptide of interest. Kits for generating and screening phage display libraries are commercially available (e.g., the Pharmacia *Recombinant Phage Antibody System*, Catalog No. 27-9400-01; and the Stratagene *SurfZAP ϕ Phage Display Kit*, Catalog No. 240612). Additionally, examples of methods and reagents particularly amenable for use in generating and screening antibody display library can be found in, for example, U.S. Patent No. 5,223,409; PCT Publication No. WO 92/18619; PCT Publication No. WO 91/17271; PCT Publication No. WO 92/20791; PCT Publication No. WO 92/15679; PCT Publication No. WO 93/01288; PCT Publication No. WO 92/01047; PCT Publication No. WO 92/09690; PCT Publication No. WO 90/02809; Fuchs et al. (1991) *Bio/Technology* 9:1370-1372; Hay et al. (1992) *Hum. Antibod. Hybridomas* 3:81-85; Huse et al. (1989) *Science* 246:1275-1281; Griffiths et al. (1993) *EMBO J.* 12:725-734.

Additionally, recombinant antibodies, such as chimeric and humanized monoclonal antibodies, comprising both human and non-human portions, which can be made using standard recombinant DNA techniques, are within the scope of the invention. Such chimeric and humanized monoclonal antibodies can be produced by recombinant DNA techniques known in the art, for example using methods described in PCT Publication No. WO 87/02671; European Patent Application 184,187; European Patent Application 171,496; European Patent Application 173,494; PCT Publication No. WO 86/01533; U.S. Patent No. 4,816,567; European Patent Application 125,023; Better et al. (1988) *Science* 240:1041-1043; Liu et al. (1987) *Proc. Natl. Acad. Sci. USA* 84:3439-3443; Liu et al. (1987) *J. Immunol.* 139:3521-3526; Sun et al. (1987) *Proc. Natl. Acad. Sci. USA* 84:214-218; Nishimura et al. (1987) *Canc. Res.* 47:999-1005; Wood et al. (1985) *Nature* 314:446-449; and Shaw et al. (1988) *J. Natl. Cancer Inst.* 80:1553-1559; Morrison (1985) *Science* 229:1202-1207; Oi et al. (1986) *Bio/Techniques* 4:214; U.S. Patent 5,225,539; Jones et al. (1986) *Nature* 321:552-525; Verhoeyan et al. (1988) *Science* 239:1534; and Beidler et al. (1988) *J. Immunol.* 141:4053-4060.

Completely human antibodies are particularly desirable for therapeutic treatment of human patients. Such antibodies can be produced using transgenic mice which are incapable of expressing endogenous immunoglobulin heavy and light chains genes, but which can express human heavy and

light chain genes. The transgenic mice are immunized in the normal fashion with a selected antigen, e.g., all or a portion of a polypeptide of the invention. Monoclonal antibodies directed against the antigen can be obtained using conventional hybridoma technology. The human immunoglobulin transgenes harbored by the transgenic mice rearrange during B cell differentiation, and subsequently undergo class switching and somatic mutation. Thus, using such a technique, it is possible to produce therapeutically useful IgG, IgA and IgE antibodies. For an overview of this technology for producing human antibodies, see Lonberg and Huszar (1995, *Int. Rev. Immunol.* 13:65-93). For a detailed discussion of this technology for producing human antibodies and human monoclonal antibodies and protocols for producing such antibodies, see, e.g., U.S. Patent 5,625,126; U.S. Patent 5,633,425; U.S. Patent 5,569,825; U.S. Patent 5,661,016; and U.S. Patent 5,545,806. In addition, companies such as Abgenix, Inc. (Freemont, CA), can be engaged to provide human antibodies directed against a selected antigen using technology similar to that described above.

Completely human antibodies which recognize a selected epitope can be generated using a technique referred to as "guided selection." In this approach a selected non-human monoclonal antibody, e.g., a murine antibody, is used to guide the selection of a completely human antibody recognizing the same epitope. (Jespers et al. (1994) *Bio/technology* 12:899-903).

An antibody directed against a polypeptide of the invention (e.g., monoclonal antibody) can be used to isolate the polypeptide by standard techniques, such as affinity chromatography or immunoprecipitation. Moreover, such an antibody can be used to detect the protein (e.g., in a cellular lysate or cell supernatant) in order to evaluate the abundance and pattern of expression of the polypeptide. The antibodies can also be used diagnostically to monitor protein levels in tissue as part of a clinical testing procedure, e.g., to, for example, determine the efficacy of a given treatment regimen. Detection can be facilitated by coupling the antibody to a detectable substance. Examples of detectable substances include various enzymes, prosthetic groups, fluorescent materials, luminescent materials, bioluminescent materials, and radioactive materials. Examples of suitable enzymes include horseradish peroxidase, alkaline phosphatase, beta-galactosidase, or acetylcholinesterase; examples of suitable prosthetic group complexes include streptavidin/biotin and avidin/biotin; examples of suitable fluorescent materials include umbelliferone, fluorescein, fluorescein isothiocyanate, rhodamine, dichlorotriazinylamine fluorescein, dansyl chloride or phycoerythrin; an example of a luminescent material includes luminol; examples of bioluminescent materials include luciferase, luciferin, and aequorin, and examples of suitable radioactive material include ^{125}I , ^{131}I , ^{35}S or ^3H .

III. Recombinant Expression Vectors and Host Cells

Another aspect of the invention pertains to vectors, preferably expression vectors, containing a nucleic acid encoding a polypeptide of the invention (or a portion thereof). As used herein, the term "vector" refers to a nucleic acid molecule capable of transporting another nucleic acid to which it has been linked. One type of vector is a "plasmid", which refers to a circular double stranded DNA loop into which additional DNA segments can be ligated. Another type of vector is a viral vector, wherein additional DNA segments can be ligated into the viral genome. Certain vectors are capable of autonomous replication in a host cell into which they are introduced (e.g., bacterial vectors having a bacterial origin of replication and episomal mammalian vectors). Other vectors (e.g., non-episomal mammalian vectors) are integrated into the genome of a host cell upon introduction into the host cell, and thereby are replicated along with the host genome. Moreover, certain vectors, expression vectors, are capable of directing the expression of genes to which they are operably linked. In general, expression vectors of utility in recombinant DNA techniques are often in the form of plasmids (vectors). However, the invention is intended to include such other forms of expression vectors, such as viral vectors (e.g., replication defective retroviruses, adenoviruses and adeno-associated viruses), which serve equivalent functions.

The recombinant expression vectors of the invention comprise a nucleic acid of the invention in a form suitable for expression of the nucleic acid in a host cell. This means that the recombinant expression vectors include one or more regulatory sequences, selected on the basis of the host cells to be used for expression, which is operably linked to the nucleic acid sequence to be expressed.

Within a recombinant expression vector, "operably linked" is intended to mean that the nucleotide sequence of interest is linked to the regulatory sequence(s) in a manner which allows for expression of the nucleotide sequence (e.g., in an *in vitro* transcription/translation system or in a host cell when the vector is introduced into the host cell). The term "regulatory sequence" is intended to include promoters, enhancers and other expression control elements (e.g., polyadenylation signals). Such regulatory sequences are described, for example, in Goeddel, *Gene Expression Technology: Methods in Enzymology* 185, Academic Press, San Diego, CA (1990). Regulatory sequences include those which direct constitutive expression of a nucleotide sequence in many types of host cell and those which direct expression of the nucleotide sequence only in certain host cells (e.g., tissue-specific regulatory sequences). It will be appreciated by those skilled in the art that the design of the expression vector can depend on such factors as the choice of the host cell to be transformed, the level of expression of protein desired, etc. The expression vectors of the invention can be introduced into host cells to thereby produce proteins or peptides, including fusion proteins or peptides, encoded by nucleic acids as described herein.

The recombinant expression vectors of the invention can be designed for expression of a polypeptide of the invention in prokaryotic (e.g., *E. coli*) or eukaryotic cells (e.g., insect cells (using

baculovirus expression vectors), yeast cells or mammalian cells). Suitable host cells are discussed further in Goeddel, *supra*. Alternatively, the recombinant expression vector can be transcribed and translated *in vitro*, for example using T7 promoter regulatory sequences and T7 polymerase.

Expression of proteins in prokaryotes is most often carried out in *E. coli* with vectors
5 containing constitutive or inducible promoters directing the expression of either fusion or non-fusion proteins. Fusion vectors add a number of amino acids to a protein encoded therein, usually to the amino terminus of the recombinant protein. Such fusion vectors typically serve three purposes: 1) to increase expression of recombinant protein; 2) to increase the solubility of the recombinant protein; and 3) to aid in the purification of the recombinant protein by acting as a ligand in affinity
10 purification. Often, in fusion expression vectors, a proteolytic cleavage site is introduced at the junction of the fusion moiety and the recombinant protein to enable separation of the recombinant protein from the fusion moiety subsequent to purification of the fusion protein. Such enzymes, and their cognate recognition sequences, include Factor Xa, thrombin and enterokinase. Typical fusion expression vectors include pGEX (Pharmacia Biotech Inc; Smith and Johnson (1988) *Gene* 67:31-
15 40), pMAL (New England Biolabs, Beverly, MA) and pRIT5 (Pharmacia, Piscataway, NJ) which fuse glutathione S-transferase (GST), maltose E binding protein, or protein A, respectively, to the target recombinant protein.

Examples of suitable inducible non-fusion *E. coli* expression vectors include pTrc (Amann et al., (1988) *Gene* 69:301-315) and pET 11d (Studier et al., *Gene Expression Technology: Methods in*
20 *Enzymology* 185, Academic Press, San Diego, California (1990) 60-89). Target gene expression from the pTrc vector relies on host RNA polymerase transcription from a hybrid trp-lac fusion promoter. Target gene expression from the pET 11d vector relies on transcription from a T7 gn10-lac fusion promoter mediated by a coexpressed viral RNA polymerase (T7 gn1). This viral polymerase is supplied by host strains BL21(DE3) or HMS174(DE3) from a resident λ prophage
25 harboring a T7 gn1 gene under the transcriptional control of the lacUV 5 promoter.

One strategy to maximize recombinant protein expression in *E. coli* is to express the protein in a host bacteria with an impaired capacity to proteolytically cleave the recombinant protein (Gottesman, *Gene Expression Technology: Methods in Enzymology* 185, Academic Press, San Diego, California (1990) 119-128). Another strategy is to alter the nucleic acid sequence of the nucleic acid
30 to be inserted into an expression vector so that the individual codons for each amino acid are those preferentially utilized in *E. coli* (Wada et al. (1992) *Nucleic Acids Res.* 20:2111-2118). Such alteration of nucleic acid sequences of the invention can be carried out by standard DNA synthesis techniques.

In another embodiment, the expression vector is a yeast expression vector. Examples of
35 vectors for expression in yeast *S. cerevisiae* include pYepSec1 (Baldari et al. (1987) *EMBO J.* 6:229-

234), pMFa (Kurjan and Herskowitz, (1982) *Cell* 30:933-943), pJRY88 (Schultz et al. (1987) *Gene* 54:113-123), pYES2 (Invitrogen Corporation, San Diego, CA), and pPicZ (Invitrogen Corp, San Diego, CA).

Alternatively, the expression vector is a baculovirus expression vector. Baculovirus vectors
5 available for expression of proteins in cultured insect cells (e.g., Sf 9 cells) include the pAc series (Smith et al. (1983) *Mol. Cell Biol.* 3:2156-2165) and the pVL series (Lucklow and Summers (1989) *Virology* 170:31-39).

In yet another embodiment, a nucleic acid of the invention is expressed in mammalian cells using a mammalian expression vector. Examples of mammalian expression vectors include pCDM8
10 (Seed (1987) *Nature* 329:840) and pMT2PC (Kaufman et al. (1987) *EMBO J.* 6:187-195). When used in mammalian cells, the expression vector's control functions are often provided by viral regulatory elements. For example, commonly used promoters are derived from polyoma, Adenovirus 2, cytomegalovirus and Simian Virus 40. For other suitable expression systems for both prokaryotic and eukaryotic cells see chapters 16 and 17 of Sambrook et al., *supra*.

In another embodiment, the recombinant mammalian expression vector is capable of
15 directing expression of the nucleic acid preferentially in a particular cell type (e.g., tissue-specific regulatory elements are used to express the nucleic acid). Tissue-specific regulatory elements are known in the art. Non-limiting examples of suitable tissue-specific promoters include the albumin promoter (liver-specific; Pinkert et al. (1987) *Genes Dev.* 1:268-277), lymphoid-specific promoters
20 (Calame and Eaton (1988) *Adv. Immunol.* 43:235-275), in particular promoters of T cell receptors (Winoto and Baltimore (1989) *EMBO J.* 8:729-733) and immunoglobulins (Banerji et al. (1983) *Cell* 33:729-740; Queen and Baltimore (1983) *Cell* 33:741-748), neuron-specific promoters (e.g., the neurofilament promoter; Byrne and Ruddle (1989) *Proc. Natl. Acad. Sci. USA* 86:5473-5477), pancreas-specific promoters (Edlund et al. (1985) *Science* 230:912-916), and mammary gland-
25 specific promoters (e.g., milk whey promoter; U.S. Patent No. 4,873,316 and European Application Publication No. 264,166). Developmentally-regulated promoters are also encompassed, for example the murine hox promoters (Kessel and Gruss (1990) *Science* 249:374-379) and the beta-fetoprotein promoter (Campes and Tilghman (1989) *Genes Dev.* 3:537-546).

The invention further provides a recombinant expression vector comprising a DNA molecule
30 of the invention cloned into the expression vector in an antisense orientation. That is, the DNA molecule is operably linked to a regulatory sequence in a manner which allows for expression (by transcription of the DNA molecule) of an RNA molecule which is antisense to the mRNA encoding a polypeptide of the invention. Regulatory sequences operably linked to a nucleic acid cloned in the antisense orientation can be chosen which direct the continuous expression of the antisense RNA
35 molecule in a variety of cell types, for instance viral promoters and/or enhancers, or regulatory

sequences can be chosen which direct constitutive, tissue specific or cell type specific expression of antisense RNA. The antisense expression vector can be in the form of a recombinant plasmid, phagemid or attenuated virus in which antisense nucleic acids are produced under the control of a high efficiency regulatory region, the activity of which can be determined by the cell type into which the vector is introduced. For a discussion of the regulation of gene expression using antisense genes see Weintraub et al. (*Reviews - Trends in Genetics*, Vol. 1(1) 1986).

Another aspect of the invention pertains to host cells into which a recombinant expression vector of the invention has been introduced. The terms "host cell" and "recombinant host cell" are used interchangeably herein. It is understood that such terms refer not only to the particular subject cell but to the progeny or potential progeny of such a cell. Because certain modifications may occur in succeeding generations due to either mutation or environmental influences, such progeny may not, in fact, be identical to the parent cell, but are still included within the scope of the term as used herein.

A host cell can be any prokaryotic (e.g., *E. coli*) or eukaryotic cell (e.g., insect cells, yeast or mammalian cells).

Vector DNA can be introduced into prokaryotic or eukaryotic cells via conventional transformation or transfection techniques. As used herein, the terms "transformation" and "transfection" are intended to refer to a variety of art-recognized techniques for introducing foreign nucleic acid into a host cell, including calcium phosphate or calcium chloride co-precipitation, DEAE-dextran-mediated transfection, lipofection, or electroporation. Suitable methods for transforming or transfecting host cells can be found in Sambrook, et al. (*supra*), and other laboratory manuals.

For stable transfection of mammalian cells, it is known that, depending upon the expression vector and transfection technique used, only a small fraction of cells may integrate the foreign DNA into their genome. In order to identify and select these integrants, a gene that encodes a selectable marker (e.g., for resistance to antibiotics) is generally introduced into the host cells along with the gene of interest. Preferred selectable markers include those which confer resistance to drugs, such as G418, hygromycin and methotrexate. Cells stably transfected with the introduced nucleic acid can be identified by drug selection (e.g., cells that have incorporated the selectable marker gene will survive, while the other cells die).

A host cell of the invention, such as a prokaryotic or eukaryotic host cell in culture, can be used to produce a polypeptide of the invention. Accordingly, the invention further provides methods for producing a polypeptide of the invention using the host cells of the invention. In one embodiment, the method comprises culturing the host cell of invention (into which a recombinant expression vector encoding a polypeptide of the invention has been introduced) in a suitable medium

such that the polypeptide is produced. In another embodiment, the method further comprises isolating the polypeptide from the medium or the host cell.

The host cells of the invention can also be used to produce nonhuman transgenic animals. For example, in one embodiment, a host cell of the invention is a fertilized oocyte or an embryonic stem cell into which a sequence encoding a polypeptide of the invention has been introduced. Such host cells can then be used to create non-human transgenic animals in which exogenous sequences encoding a polypeptide of the invention have been introduced into their genome or homologous recombinant animals in which endogenous encoding a polypeptide of the invention sequences have been altered. Such animals are useful for studying the function and/or activity of the polypeptide and for identifying and/or evaluating modulators of polypeptide activity. As used herein, a "transgenic animal" is a non-human animal, preferably a mammal, more preferably a rodent such as a rat or mouse, in which one or more of the cells of the animal includes a transgene. Other examples of transgenic animals include non-human primates, sheep, dogs, cows, goats, chickens, amphibians, etc.

A transgene is exogenous DNA which is integrated into the genome of a cell from which a transgenic animal develops and which remains in the genome of the mature animal, thereby directing the expression of an encoded gene product in one or more cell types or tissues of the transgenic animal. As used herein, an "homologous recombinant animal" is a non-human animal, preferably a mammal, more preferably a mouse, in which an endogenous gene has been altered by homologous recombination between the endogenous gene and an exogenous DNA molecule introduced into a cell of the animal, e.g., an embryonic cell of the animal, prior to development of the animal.

A transgenic animal of the invention can be created by introducing nucleic acid encoding a polypeptide of the invention (or a homologue thereof) into the male pronuclei of a fertilized oocyte, e.g., by microinjection, retroviral infection, and allowing the oocyte to develop in a pseudopregnant female foster animal. Intronic sequences and polyadenylation signals can also be included in the transgene to increase the efficiency of expression of the transgene. A tissue-specific regulatory sequence(s) can be operably linked to the transgene to direct expression of the polypeptide of the invention to particular cells. Methods for generating transgenic animals via embryo manipulation and microinjection, particularly animals such as mice, have become conventional in the art and are described, for example, in U.S. Patent Nos. 4,736,866 and 4,870,009, U.S. Patent No. 4,873,191 and in Hogan, *Manipulating the Mouse Embryo*, (Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., 1986). Similar methods are used for production of other transgenic animals. A transgenic founder animal can be identified based upon the presence of the transgene in its genome and/or expression of mRNA encoding the transgene in tissues or cells of the animals. A transgenic founder animal can then be used to breed additional animals carrying the transgene. Moreover,

transgenic animals carrying the transgene can further be bred to other transgenic animals carrying other transgenes.

To create an homologous recombinant animal, a vector is prepared which contains at least a portion of a gene encoding a polypeptide of the invention into which a deletion, addition or substitution has been introduced to thereby alter, e.g., functionally disrupt, the gene. In a preferred embodiment, the vector is designed such that, upon homologous recombination, the endogenous gene is functionally disrupted (i.e., no longer encodes a functional protein; also referred to as a "knock out" vector). Alternatively, the vector can be designed such that, upon homologous recombination, the endogenous gene is mutated or otherwise altered but still encodes functional protein (e.g., the upstream regulatory region can be altered to thereby alter the expression of the endogenous protein). In the homologous recombination vector, the altered portion of the gene is flanked at its 5' and 3' ends by additional nucleic acid of the gene to allow for homologous recombination to occur between the exogenous gene carried by the vector and an endogenous gene in an embryonic stem cell. The additional flanking nucleic acid sequences are of sufficient length for successful homologous recombination with the endogenous gene. Typically, several kilobases of flanking DNA (both at the 5' and 3' ends) are included in the vector (*see, e.g.,* Thomas and Capecchi (1987) *Cell* 51:503 for a description of homologous recombination vectors). The vector is introduced into an embryonic stem cell line (e.g., by electroporation) and cells in which the introduced gene has homologously recombined with the endogenous gene are selected (*see, e.g.,* Li et al. (1992) *Cell* 69:915). The selected cells are then injected into a blastocyst of an animal (e.g., a mouse) to form aggregation chimeras (*see, e.g.,* Bradley in *Teratocarcinomas and Embryonic Stem Cells: A Practical Approach*, Robertson, ed. (IRL, Oxford, 1987) pp. 113-152). A chimeric embryo can then be implanted into a suitable pseudopregnant female foster animal and the embryo brought to term. Progeny harboring the homologously recombined DNA in their germ cells can be used to breed animals in which all cells of the animal contain the homologously recombined DNA by germline transmission of the transgene. Methods for constructing homologous recombination vectors and homologous recombinant animals are described further in Bradley (1991) *Current Opinion in Bio/Technology* 2:823-829 and in PCT Publication Nos. WO 90/11354, WO 91/01140, WO 92/0968, and WO 93/04169.

In another embodiment, transgenic non-human animals can be produced which contain selected systems which allow for regulated expression of the transgene. One example of such a system is the *cre/loxP* recombinase system of bacteriophage P1. For a description of the *cre/loxP* recombinase system, *see, e.g.,* Lakso et al. (1992) *Proc. Natl. Acad. Sci. USA* 89:6232-6236. Another example of a recombinase system is the FLP recombinase system of *Saccharomyces cerevisiae* (O'Gorman et al. (1991) *Science* 251:1351-1355. If a *cre/loxP* recombinase system is used

to regulate expression of the transgene, animals containing transgenes encoding both the *Cre* recombinase and a selected protein are required. Such animals can be provided through the construction of "double" transgenic animals, e.g., by mating two transgenic animals, one containing a transgene encoding a selected protein and the other containing a transgene encoding a recombinase.

5 Clones of the non-human transgenic animals described herein can also be produced according to the methods described in Wilmut et al. (1997) *Nature* 385:810-813 and PCT Publication NOS. WO 97/07668 and WO 97/07669.

IV. Pharmaceutical Compositions

10 The nucleic acid molecules, polypeptides, and antibodies (also referred to herein as "active compounds") of the invention can be incorporated into pharmaceutical compositions suitable for administration. Such compositions typically comprise the nucleic acid molecule, protein, or antibody and a pharmaceutically acceptable carrier. As used herein the language "pharmaceutically acceptable carrier" is intended to include any and all solvents, dispersion media, coatings, antibacterial and
15 antifungal agents, isotonic and absorption delaying agents, and the like, compatible with pharmaceutical administration. The use of such media and agents for pharmaceutically active substances is well known in the art. Except insofar as any conventional media or agent is incompatible with the active compound, use thereof in the compositions is contemplated. Supplementary active compounds can also be incorporated into the compositions.

20 The invention includes methods for preparing pharmaceutical compositions for modulating the expression or activity of a polypeptide or nucleic acid of the invention. Such methods comprise formulating a pharmaceutically acceptable carrier with an agent which modulates expression or activity of a polypeptide or nucleic acid of the invention. Such compositions can further include additional active agents. Thus, the invention further includes methods for preparing a pharmaceutical
25 composition by formulating a pharmaceutically acceptable carrier with an agent which modulates expression or activity of a polypeptide or nucleic acid of the invention and one or more additional active compounds.

A pharmaceutical composition of the invention is formulated to be compatible with its intended route of administration. Examples of routes of administration include parenteral, e.g.,
30 intravenous, intradermal, subcutaneous, oral (e.g., inhalation), transdermal (topical), transmucosal, and rectal administration. Solutions or suspensions used for parenteral, intradermal, or subcutaneous application can include the following components: a sterile diluent such as water for injection, saline solution, fixed oils, polyethylene glycols, glycerine, propylene glycol or other synthetic solvents; antibacterial agents such as benzyl alcohol or methyl parabens; antioxidants such as ascorbic acid or
35 sodium bisulfite; chelating agents such as ethylenediaminetetraacetic acid; buffers such as acetates,

citrates or phosphates and agents for the adjustment of tonicity such as sodium chloride or dextrose. pH can be adjusted with acids or bases, such as hydrochloric acid or sodium hydroxide. The parenteral preparation can be enclosed in ampoules, disposable syringes or multiple dose vials made of glass or plastic.

5 Pharmaceutical compositions suitable for injectable use include sterile aqueous solutions (where water soluble) or dispersions and sterile powders for the extemporaneous preparation of sterile injectable solutions or dispersions. For intravenous administration, suitable carriers include physiological saline, bacteriostatic water, Cremophor EL® (BASF; Parsippany, NJ) or phosphate buffered saline (PBS). In all cases, the composition must be sterile and should be fluid to the extent
10 that easy syringability exists. It must be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms such as bacteria and fungi. The carrier can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (for example, glycerol, propylene glycol, and liquid polyethylene glycol, and the like), and suitable mixtures thereof. The proper fluidity can be maintained, for example, by the use of a coating such as
15 lecithin, by the maintenance of the required particle size in the case of dispersion and by the use of surfactants. Prevention of the action of microorganisms can be achieved by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, ascorbic acid, thimerosal, and the like. In many cases, it will be preferable to include isotonic agents, for example, sugars, polyalcohols such as mannitol, sorbitol, sodium chloride in the composition. Prolonged absorption of the
20 injectable compositions can be brought about by including in the composition an agent which delays absorption, for example, aluminum monostearate and gelatin.

 Sterile injectable solutions can be prepared by incorporating the active compound (e.g., a polypeptide or antibody) in the required amount in an appropriate solvent with one or a combination of ingredients enumerated above, as required, followed by filtered sterilization. Generally,
25 dispersions are prepared by incorporating the active compound into a sterile vehicle which contains a basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum drying and freeze-drying which yields a powder of the active ingredient plus any additional desired ingredient from a previously sterile-filtered solution thereof.

30 Oral compositions generally include an inert diluent or an edible carrier. They can be enclosed in gelatin capsules or compressed into tablets. For the purpose of oral therapeutic administration, the active compound can be incorporated with excipients and used in the form of tablets, troches, or capsules. Oral compositions can also be prepared using a fluid carrier for use as a mouthwash, wherein the compound in the fluid carrier is applied orally and swished and expectorated
35 or swallowed.

Pharmaceutically compatible binding agents, and/or adjuvant materials can be included as part of the composition. The tablets, pills, capsules, troches and the like can contain any of the following ingredients, or compounds of a similar nature: a binder such as microcrystalline cellulose, gum tragacanth or gelatin; an excipient such as starch or lactose, a disintegrating agent such as alginic acid, Primogel, or corn starch; a lubricant such as magnesium stearate or Sterotes; a glidant such as colloidal silicon dioxide; a sweetening agent such as sucrose or saccharin; or a flavoring agent such as peppermint, methyl salicylate, or orange flavoring.

For administration by inhalation, the compounds are delivered in the form of an aerosol spray from a pressurized container or dispenser which contains a suitable propellant, e.g., a gas such as carbon dioxide, or a nebulizer.

Systemic administration can also be by transmucosal or transdermal means. For transmucosal or transdermal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art, and include, for example, for transmucosal administration, detergents, bile salts, and fusidic acid derivatives. Transmucosal administration can be accomplished through the use of nasal sprays or suppositories. For transdermal administration, the active compounds are formulated into ointments, salves, gels, or creams as generally known in the art.

The compounds can also be prepared in the form of suppositories (e.g., with conventional suppository bases such as cocoa butter and other glycerides) or retention enemas for rectal delivery.

In one embodiment, the active compounds are prepared with carriers that will protect the compound against rapid elimination from the body, such as a controlled release formulation, including implants and microencapsulated delivery systems. Biodegradable, biocompatible polymers can be used, such as ethylene vinyl acetate, polyanhydrides, polyglycolic acid, collagen, polyorthoesters, and polylactic acid. Methods for preparation of such formulations will be apparent to those skilled in the art. The materials can also be obtained commercially from Alza Corporation and Nova Pharmaceuticals, Inc. Liposomal suspensions (including liposomes targeted to infected cells with monoclonal antibodies to viral antigens) can also be used as pharmaceutically acceptable carriers. These can be prepared according to methods known to those skilled in the art, for example, as described in U.S. Patent No. 4,522,811.

It is especially advantageous to formulate oral or parenteral compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used herein refers to physically discrete units suited as unitary dosages for the subject to be treated; each unit containing a predetermined quantity of active compound calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. The specification for the dosage unit forms of the invention are dictated by and directly dependent on the unique characteristics of the active

compound and the particular therapeutic effect to be achieved, and the limitations inherent in the art of compounding such an active compound for the treatment of individuals.

As defined herein, a therapeutically effective amount of antibody, protein, or polypeptide (i.e., an effective dosage) ranges from about 0.001 to about 30, 40, 50, 60, 70, 80, 90, 100 mg/kg of body weight, about 0.001 to 30 mg/kg body weight, preferably about 0.01 to 25 mg/kg body weight, more preferably about 0.1 to 20 mg/kg (e.g., generally about 10 mg/kg to 20 mg/kg body weight) body weight, and even more preferably about 1 to 10 mg/kg, 2 to 9 mg/kg, 3 to 8 mg/kg, 4 to 7 mg/kg, or 5 to 6 mg/kg body weight.

With regard to antibodies which are to act in the brain, a dosage of 50 mg/kg to 100 mg/kg is typically appropriate. Generally, partially human antibodies and fully human antibodies have a longer half-life within the human body than other antibodies. Accordingly, lower dosages and less frequent administration is often possible. Modifications such as lipidation can be used to stabilize antibodies and to enhance uptake and tissue penetration (e.g., into the brain). A method for lipidation of antibodies is described by Cruikshank et al. ((1997) *J. Acquired Immune Deficiency Syndromes and Human Retrovirology* 14:193).

The skilled artisan will appreciate that certain factors may influence the dosage required to effectively treat a subject, including but not limited to the severity of the disease or disorder, previous treatments, the general health and/or age of the subject, and other diseases present. Moreover, treatment of a subject with a therapeutically effective amount of a protein, polypeptide, or antibody can include a single treatment or, preferably, can include a series of treatments. In one embodiment, a subject is treated with a protein, or polypeptide in the range of between about 0.1 to 20 mg/kg body weight, one time per week for between about 1 to 10 weeks, preferably between about 2 to 8 weeks, more preferably between about 3 to 8 weeks, and even more preferably for about 4, 5, or 6 weeks. It will also be appreciated that the effective dosage of antibody, protein, or polypeptide used for treatment may increase or decrease over the course of a particular treatment. Changes in dosage may result and become apparent from the results of diagnostic assays as described herein.

The nucleic acid molecules of the invention can be inserted into vectors and used as gene therapy vectors. Gene therapy vectors can be delivered to a subject by, for example, intravenous injection, local administration (U.S. Patent 5,328,470) or by stereotactic injection (*see, e.g.*, Chen et al. (1994) *Proc. Natl. Acad. Sci. USA* 91:3054-3057). The pharmaceutical preparation of the gene therapy vector can include the gene therapy vector in an acceptable diluent, or can comprise a slow release matrix in which the gene delivery vehicle is imbedded. Alternatively, where the complete gene delivery vector can be produced intact from recombinant cells, e.g., retroviral vectors, the pharmaceutical preparation can include one or more cells which produce the gene delivery system.

The pharmaceutical compositions can be included in a container, pack, or dispenser together with instructions for administration.

V. Uses and Methods of the Invention

5 The nucleic acid molecules, proteins, protein homologues, and antibodies described herein can be used in one or more of the following methods: a) screening assays; b) detection assays (e.g., chromosomal mapping, tissue typing, forensic biology); c) predictive medicine (e.g., diagnostic assays, prognostic assays, monitoring clinical trials, and pharmacogenomics); and d) methods of treatment (e.g., therapeutic and prophylactic). The isolated nucleic acid molecules of the invention
10 can be used to express proteins (e.g., via a recombinant expression vector in a host cell in gene therapy applications), to detect mRNA (e.g., in a biological sample) or a genetic lesion, and to modulate activity of a polypeptide of the invention. In addition, the polypeptides of the invention can be used to screen drugs or compounds which modulate activity or expression of a polypeptide of the invention as well as to treat disorders characterized by insufficient or excessive production of a
15 protein of the invention or production of a form of a protein of the invention which has decreased or aberrant activity compared to the wild type protein. In addition, the antibodies of the invention can be used to detect and isolate a protein of the and modulate activity of a protein of the invention.

 This invention further pertains to novel agents identified by the above-described screening assays and uses thereof for treatments as described herein.

A. Screening Assays

 The invention provides a method (also referred to herein as a "screening assay") for identifying modulators, i.e., candidate or test compounds or agents (e.g., peptides, peptidomimetics, small molecules or other drugs) which bind to polypeptide of the invention or have a stimulatory or
25 inhibitory effect on, for example, expression or activity of a polypeptide of the invention.

 In one embodiment, the invention provides assays for screening candidate or test compounds which bind to or modulate the activity of the membrane-bound form of a polypeptide of the invention or biologically active portion thereof. The test compounds of the present invention can be obtained using any of the numerous approaches in combinatorial library methods known in the art, including:
30 biological libraries; spatially addressable parallel solid phase or solution phase libraries; synthetic library methods requiring deconvolution; the "one-bead one-compound" library method; and synthetic library methods using affinity chromatography selection. The biological library approach is limited to peptide libraries, while the other four approaches are applicable to peptide, non-peptide oligomer or small molecule libraries of compounds (Lam (1997) *Anticancer Drug Des.* 12:145).

Examples of methods for the synthesis of molecular libraries can be found in the art, for example in: DeWitt et al. (1993) *Proc. Natl. Acad. Sci. USA* 90:6909; Erb et al. (1994) *Proc. Natl. Acad. Sci. USA* 91:11422; Zuckermann et al. (1994). *J. Med. Chem.* 37:2678; Cho et al. (1993) *Science* 261:1303; Carrell et al. (1994) *Angew. Chem. Int. Ed. Engl.* 33:2059; Carell et al. (1994) *Angew. Chem. Int. Ed. Engl.* 33:2061; and Gallop et al. (1994) *J. Med. Chem.* 37:1233.

Libraries of compounds may be presented in solution (e.g., Houghten (1992) *Bio/Techniques* 13:412-421), or on beads (Lam (1991) *Nature* 354:82-84), chips (Fodor (1993) *Nature* 364:555-556), bacteria (U.S. Patent No. 5,223,409), spores (Patent NOS. 5,571,698; 5,403,484; and 5,223,409), plasmids (Cull et al. (1992) *Proc. Natl. Acad. Sci. USA* 89:1865-1869) or phage (Scott and Smith (1990) *Science* 249:386-390; Devlin (1990) *Science* 249:404-406; Cwirla et al. (1990) *Proc. Natl. Acad. Sci. USA* 87:6378-6382; and Felici (1991) *J. Mol. Biol.* 222:301-310).

In one embodiment, an assay is a cell-based assay in which a cell which expresses a membrane-bound form of a polypeptide of the invention, or a biologically active portion thereof, on the cell surface is contacted with a test compound and the ability of the test compound to bind to the polypeptide determined. The cell, for example, can be a yeast cell or a cell of mammalian origin. Determining the ability of the test compound to bind to the polypeptide can be accomplished, for example, by coupling the test compound with a radioisotope or enzymatic label such that binding of the test compound to the polypeptide or biologically active portion thereof can be determined by detecting the labeled compound in a complex. For example, test compounds can be labeled with ¹²⁵I, ³⁵S, ¹⁴C, or ³H, either directly or indirectly, and the radioisotope detected by direct counting of radioemmission or by scintillation counting. Alternatively, test compounds can be enzymatically labeled with, for example, horseradish peroxidase, alkaline phosphatase, or luciferase, and the enzymatic label detected by determination of conversion of an appropriate substrate to product. In a preferred embodiment, the assay comprises contacting a cell which expresses a membrane-bound form of a polypeptide of the invention, or a biologically active portion thereof, on the cell surface with a known compound which binds the polypeptide to form an assay mixture, contacting the assay mixture with a test compound, and determining the ability of the test compound to interact with the polypeptide, wherein determining the ability of the test compound to interact with the polypeptide comprises determining the ability of the test compound to preferentially bind to the polypeptide or a biologically active portion thereof as compared to the known compound.

In another embodiment, an assay is a cell-based assay comprising contacting a cell expressing a membrane-bound form of a polypeptide of the invention, or a biologically active portion thereof, on the cell surface with a test compound and determining the ability of the test compound to modulate (e.g., stimulate or inhibit) the activity of the polypeptide or biologically active portion thereof. Determining the ability of the test compound to modulate the activity of the polypeptide or a

biologically active portion thereof can be accomplished, for example, by determining the ability of the polypeptide protein to bind to or interact with a target molecule.

Determining the ability of a polypeptide of the invention to bind to or interact with a target molecule can be accomplished by one of the methods described above for determining direct binding.

5 As used herein, a "target molecule" is a molecule with which a selected polypeptide (e.g., a polypeptide of the invention) binds or interacts with in nature, for example, a molecule on the surface of a cell which expresses the selected protein, a molecule on the surface of a second cell, a molecule in the extracellular milieu, a molecule associated with the internal surface of a cell membrane or a cytoplasmic molecule. A target molecule can be a polypeptide of the invention or some other
10 polypeptide or protein. For example, a target molecule can be a component of a signal transduction pathway which facilitates transduction of an extracellular signal (e.g., a signal generated by binding of a compound to a polypeptide of the invention) through the cell membrane and into the cell or a second intercellular protein which has catalytic activity or a protein which facilitates the association of downstream signaling molecules with a polypeptide of the invention. Determining the ability of a
15 polypeptide of the invention to bind to or interact with a target molecule can be accomplished by determining the activity of the target molecule. For example, the activity of the target molecule can be determined by detecting induction of a cellular second messenger of the target (e.g., intracellular Ca^{2+} , diacylglycerol, IP3, etc.), detecting catalytic/enzymatic activity of the target on an appropriate substrate, detecting the induction of a reporter gene (e.g., a regulatory element that is responsive to a
20 polypeptide of the invention operably linked to a nucleic acid encoding a detectable marker, e.g., luciferase), or detecting a cellular response, for example, cellular differentiation, or cell proliferation.

In yet another embodiment, an assay of the present invention is a cell-free assay comprising contacting a polypeptide of the invention or biologically active portion thereof with a test compound and determining the ability of the test compound to bind to the polypeptide or biologically active
25 portion thereof. Binding of the test compound to the polypeptide can be determined either directly or indirectly as described above. In a preferred embodiment, the assay includes contacting the polypeptide of the invention or biologically active portion thereof with a known compound which binds the polypeptide to form an assay mixture, contacting the assay mixture with a test compound, and determining the ability of the test compound to interact with the polypeptide, wherein
30 determining the ability of the test compound to interact with the polypeptide comprises determining the ability of the test compound to preferentially bind to the polypeptide or biologically active portion thereof as compared to the known compound.

In another embodiment, an assay is a cell-free assay comprising contacting a polypeptide of the invention or biologically active portion thereof with a test compound and determining the ability
35 of the test compound to modulate (e.g., stimulate or inhibit) the activity of the polypeptide or

biologically active portion thereof. Determining the ability of the test compound to modulate the activity of the polypeptide can be accomplished, for example, by determining the ability of the polypeptide to bind to a target molecule by one of the methods described above for determining direct binding. In an alternative embodiment, determining the ability of the test compound to modulate the activity of the polypeptide can be accomplished by determining the ability of the polypeptide of the invention to further modulate the target molecule. For example, the catalytic/enzymatic activity of the target molecule on an appropriate substrate can be determined as previously described.

In yet another embodiment, the cell-free assay comprises contacting a polypeptide of the invention or biologically active portion thereof with a known compound which binds the polypeptide to form an assay mixture, contacting the assay mixture with a test compound, and determining the ability of the test compound to interact with the polypeptide, wherein determining the ability of the test compound to interact with the polypeptide comprises determining the ability of the polypeptide to preferentially bind to or modulate the activity of a target molecule.

The cell-free assays of the present invention are amenable to use of both a soluble form or the membrane-bound form of a polypeptide of the invention. In the case of cell-free assays comprising the membrane-bound form of the polypeptide, it may be desirable to utilize a solubilizing agent such that the membrane-bound form of the polypeptide is maintained in solution. Examples of such solubilizing agents include non-ionic detergents such as n-octylglucoside, n-dodecylglucoside, n-octylmaltoside, octanoyl-predicted N-methylglucamide, decanoyl-predicted N-methylglucamide, Triton X-100, Triton X-114, Thesit, Isotridecypoly(ethylene glycol ether)_n, 3-[(3-cholamidopropyl)dimethylamminio]-1-propane sulfonate (CHAPS), 3-[(3-cholamidopropyl)dimethylamminio]-2-hydroxy-1-propane sulfonate (CHAPSO), or predicted N-dodecyl=N,predicted N-dimethyl-3-ammonio-1-propane sulfonate.

In more than one embodiment of the above assay methods of the present invention, it may be desirable to immobilize either the polypeptide of the invention or its target molecule to facilitate separation of complexed from uncomplexed forms of one or both of the proteins, as well as to accommodate automation of the assay. Binding of a test compound to the polypeptide, or interaction of the polypeptide with a target molecule in the presence and absence of a candidate compound, can be accomplished in any vessel suitable for containing the reactants. Examples of such vessels include microtitre plates, test tubes, and micro-centrifuge tubes. In one embodiment, a fusion protein can be provided which adds a domain that allows one or both of the proteins to be bound to a matrix. For example, glutathione-S-transferase fusion proteins or glutathione-S-transferase fusion proteins can be adsorbed onto glutathione sepharose beads (Sigma Chemical; St. Louis, MO) or glutathione derivatized microtitre plates, which are then combined with the test compound or the test compound

and either the non-adsorbed target protein or A polypeptide of the invention, and the mixture incubated under conditions conducive to complex formation (e.g., at physiological conditions for salt and pH). Following incubation, the beads or microtitre plate wells are washed to remove any unbound components and complex formation is measured either directly or indirectly, for example, as described above. Alternatively, the complexes can be dissociated from the matrix, and the level of binding or activity of the polypeptide of the invention can be determined using standard techniques.

Other techniques for immobilizing proteins on matrices can also be used in the screening assays of the invention. For example, either the polypeptide of the invention or its target molecule can be immobilized utilizing conjugation of biotin and streptavidin. Biotinylated polypeptide of the invention or target molecules can be prepared from biotin-NHS (predicted N-hydroxy-succinimide) using techniques well known in the art (e.g., biotinylation kit, Pierce Chemicals; Rockford, IL), and immobilized in the wells of streptavidin-coated 96 well plates (Pierce Chemical). Alternatively, antibodies reactive with the polypeptide of the invention or target molecules but which do not interfere with binding of the polypeptide of the invention to its target molecule can be derivatized to the wells of the plate, and unbound target or polypeptide of the invention trapped in the wells by antibody conjugation. Methods for detecting such complexes, in addition to those described above for the GST-immobilized complexes, include immunodetection of complexes using antibodies reactive with the polypeptide of the invention or target molecule, as well as enzyme-linked assays which rely on detecting an enzymatic activity associated with the polypeptide of the invention or target molecule.

In another embodiment, modulators of expression of a polypeptide of the invention are identified in a method in which a cell is contacted with a candidate compound and the expression of the selected mRNA or protein (i.e., the mRNA or protein corresponding to a polypeptide or nucleic acid of the invention) in the cell is determined. The level of expression of the selected mRNA or protein in the presence of the candidate compound is compared to the level of expression of the selected mRNA or protein in the absence of the candidate compound. The candidate compound can then be identified as a modulator of expression of the polypeptide of the invention based on this comparison. For example, when expression of the selected mRNA or protein is greater (statistically significantly greater) in the presence of the candidate compound than in its absence, the candidate compound is identified as a stimulator of the selected mRNA or protein expression. Alternatively, when expression of the selected mRNA or protein is less (statistically significantly less) in the presence of the candidate compound than in its absence, the candidate compound is identified as an inhibitor of the selected mRNA or protein expression. The level of the selected mRNA or protein expression in the cells can be determined by methods described herein.

In yet another aspect of the invention, a polypeptide of the inventions can be used as "bait proteins" in a two-hybrid assay or three hybrid assay (*see, e.g.*, U.S. Patent No. 5,283,317; Zervos et al. (1993) *Cell* 72:223-232; Madura et al. (1993) *J. Biol. Chem.* 268:12046-12054; Bartel et al. (1993) *Bio/Techniques* 14:920-924; Iwabuchi et al. (1993) *Oncogene* 8:1693-1696; and PCT Publication No. WO 94/10300), to identify other proteins, which bind to or interact with the polypeptide of the invention and modulate activity of the polypeptide of the invention. Such binding proteins are also likely to be involved in the propagation of signals by the polypeptide of the inventions as, for example, upstream or downstream elements of a signaling pathway involving the polypeptide of the invention.

This invention further pertains to novel agents identified by the above-described screening assays and uses thereof for treatments as described herein.

B. Detection Assays

Portions or fragments of the cDNA sequences identified herein (and the corresponding complete gene sequences) can be used in numerous ways as polynucleotide reagents. For example, these sequences can be used to: (i) map their respective genes on a chromosome and, thus, locate gene regions associated with genetic disease; (ii) identify an individual from a minute biological sample (tissue typing); and (iii) aid in forensic identification of a biological sample. These applications are described in the subsections below.

1. Chromosome Mapping

Once the sequence (or a portion of the sequence) of a gene has been isolated, this sequence can be used to map the location of the gene on a chromosome. Accordingly, nucleic acid molecules described herein or fragments thereof, can be used to map the location of the corresponding genes on a chromosome. The mapping of the sequences to chromosomes is an important first step in correlating these sequences with genes associated with disease.

Briefly, genes can be mapped to chromosomes by preparing PCR primers (preferably 15-25 bp in length) from the sequence of a gene of the invention. Computer analysis of the sequence of a gene of the invention can be used to rapidly select primers that do not span more than one exon in the genomic DNA, thus complicating the amplification process. These primers can then be used for PCR screening of somatic cell hybrids containing individual human chromosomes. Only those hybrids containing the human gene corresponding to the gene sequences will yield an amplified fragment. For a review of this technique, see D'Eustachio et al. ((1983) *Science* 220:919-924).

PCR mapping of somatic cell hybrids is a rapid procedure for assigning a particular sequence to a particular chromosome. Three or more sequences can be assigned per day using a single thermal

cycler. Using the nucleic acid sequences of the invention to design oligonucleotide primers, sublocalization can be achieved with panels of fragments from specific chromosomes. Other mapping strategies which can similarly be used to map a gene to its chromosome include *in situ* hybridization (described in Fan et al. (1990) *Proc. Natl. Acad. Sci. USA* 87:6223-27), pre-screening
5 with labeled flow-sorted chromosomes (CITE), and pre-selection by hybridization to chromosome specific cDNA libraries. Fluorescence *in situ* hybridization (FISH) of a DNA sequence to a metaphase chromosomal spread can further be used to provide a precise chromosomal location in one step. For a review of this technique, see Verma et al., (Human Chromosomes: A Manual of Basic Techniques (Pergamon Press, New York, 1988)).

10 Reagents for chromosome mapping can be used individually to mark a single chromosome or a single site on that chromosome, or panels of reagents can be used for marking multiple sites and/or multiple chromosomes. Reagents corresponding to noncoding regions of the genes actually are preferred for mapping purposes. Coding sequences are more likely to be conserved within gene families, thus increasing the chance of cross hybridizations during chromosomal mapping.

15 Once a sequence has been mapped to a precise chromosomal location, the physical position of the sequence on the chromosome can be correlated with genetic map data. (Such data are found, for example, in V. McKusick, Mendelian Inheritance in Man, available on-line through Johns Hopkins University Welch Medical Library). The relationship between genes and disease, mapped to the same chromosomal region, can then be identified through linkage analysis (co-inheritance of
20 physically adjacent genes), described in, e.g., Egeland et al. (1987) *Nature* 325:783-787.

Moreover, differences in the DNA sequences between individuals affected and unaffected with a disease associated with a gene of the invention can be determined. If a mutation is observed in some or all of the affected individuals but not in any unaffected individuals, then the mutation is likely to be the causative agent of the particular disease. Comparison of affected and unaffected
25 individuals generally involves first looking for structural alterations in the chromosomes such as deletions or translocations that are visible from chromosome spreads or detectable using PCR based on that DNA sequence. Ultimately, complete sequencing of genes from several individuals can be performed to confirm the presence of a mutation and to distinguish mutations from polymorphisms.

30 2. Tissue Typing

The nucleic acid sequences of the present invention can also be used to identify individuals from minute biological samples. The United States military, for example, is considering the use of restriction fragment length polymorphism (RFLP) for identification of its personnel. In this technique, an individual's genomic DNA is digested with one or more restriction enzymes, and
35 probed on a Southern blot to yield unique bands for identification. This method does not suffer from

the current limitations of "Dog Tags" which can be lost, switched, or stolen, making positive identification difficult. The sequences of the present invention are useful as additional DNA markers for RFLP (described in U.S. Patent 5,272,057).

Furthermore, the sequences of the present invention can be used to provide an alternative technique which determines the actual base-by-base DNA sequence of selected portions of an individual's genome. Thus, the nucleic acid sequences described herein can be used to prepare two PCR primers from the 5' and 3' ends of the sequences. These primers can then be used to amplify an individual's DNA and subsequently sequence it.

Panels of corresponding DNA sequences from individuals, prepared in this manner, can provide unique individual identifications, as each individual will have a unique set of such DNA sequences due to allelic differences. The sequences of the present invention can be used to obtain such identification sequences from individuals and from tissue. The nucleic acid sequences of the invention uniquely represent portions of the human genome. Allelic variation occurs to some degree in the coding regions of these sequences, and to a greater degree in the noncoding regions. It is estimated that allelic variation between individual humans occurs with a frequency at about once per each 500 bases. Each of the sequences described herein can, to some degree, be used as a standard against which DNA from an individual can be compared for identification purposes. Because greater numbers of polymorphisms occur in the noncoding regions, fewer sequences are necessary to differentiate individuals. The noncoding sequences of SEQ ID NO:1 or 19 can comfortably provide positive individual identification with a panel of perhaps 10 to 1,000 primers which each yield a noncoding amplified sequence of 100 bases. If predicted coding sequences, such as those in SEQ ID NO:2 or 20 are used, a more appropriate number of primers for positive individual identification would be 500-2,000.

If a panel of reagents from the nucleic acid sequences described herein is used to generate a unique identification database for an individual, those same reagents can later be used to identify tissue from that individual. Using the unique identification database, positive identification of the individual, living or dead, can be made from extremely small tissue samples.

3. Use of Partial Gene Sequences in Forensic Biology

DNA-based identification techniques can also be used in forensic biology. Forensic biology is a scientific field employing genetic typing of biological evidence found at a crime scene as a means for positively identifying, for example, a perpetrator of a crime. To make such an identification, PCR technology can be used to amplify DNA sequences taken from very small biological samples such as tissues, e.g., hair or skin, or body fluids, e.g., blood, saliva, or semen

found at a crime scene. The amplified sequence can then be compared to a standard, thereby allowing identification of the origin of the biological sample.

The sequences of the present invention can be used to provide polynucleotide reagents, e.g., PCR primers, targeted to specific loci in the human genome, which can enhance the reliability of DNA-based forensic identifications by, for example, providing another "identification marker" (i.e. another DNA sequence that is unique to a particular individual). As mentioned above, actual base sequence information can be used for identification as an accurate alternative to patterns formed by restriction enzyme generated fragments. Sequences targeted to noncoding regions are particularly appropriate for this use as greater numbers of polymorphisms occur in the noncoding regions, making it easier to differentiate individuals using this technique. Examples of polynucleotide reagents include the nucleic acid sequences of the invention or portions thereof, e.g., fragments derived from noncoding regions having a length of at least 20 or 30 bases.

The nucleic acid sequences described herein can further be used to provide polynucleotide reagents, e.g., labeled or labelable probes which can be used in, for example, an *in situ* hybridization technique, to identify a specific tissue, e.g., brain tissue. This can be very useful in cases where a forensic pathologist is presented with a tissue of unknown origin. Panels of such probes can be used to identify tissue by species and/or by organ type.

C. Predictive Medicine

The present invention also pertains to the field of predictive medicine in which diagnostic assays, prognostic assays, pharmacogenomics, and monitoring clinical trials are used for prognostic (predictive) purposes to thereby treat an individual prophylactically. Accordingly, one aspect of the present invention relates to diagnostic assays for determining expression of a polypeptide or nucleic acid of the invention and/or activity of a polypeptide of the invention, in the context of a biological sample (e.g., blood, serum, cells, tissue) to thereby determine whether an individual is afflicted with a disease or disorder, or is at risk of developing a disorder, associated with aberrant expression or activity of a polypeptide of the invention. The invention also provides for prognostic (or predictive) assays for determining whether an individual is at risk of developing a disorder associated with aberrant expression or activity of a polypeptide of the invention. For example, mutations in a gene of the invention can be assayed in a biological sample. Such assays can be used for prognostic or predictive purpose to thereby prophylactically treat an individual prior to the onset of a disorder characterized by or associated with aberrant expression or activity of a polypeptide of the invention.

Another aspect of the invention provides methods for expression of a nucleic acid or polypeptide of the invention or activity of a polypeptide of the invention in an individual to thereby select appropriate therapeutic or prophylactic agents for that individual (referred to herein as

"pharmacogenomics"). Pharmacogenomics allows for the selection of agents (e.g., drugs) for therapeutic or prophylactic treatment of an individual based on the genotype of the individual (e.g., the genotype of the individual examined to determine the ability of the individual to respond to a particular agent).

Yet another aspect of the invention pertains to monitoring the influence of agents (e.g., drugs or other compounds) on the expression or activity of a polypeptide of the invention in clinical trials. These and other agents are described in further detail in the following sections.

1. Diagnostic Assays

An exemplary method for detecting the presence or absence of a polypeptide or nucleic acid of the invention in a biological sample involves obtaining a biological sample from a test subject and contacting the biological sample with a compound or an agent capable of detecting a polypeptide or nucleic acid (e.g., mRNA, genomic DNA) of the invention such that the presence of a polypeptide or nucleic acid of the invention is detected in the biological sample. A preferred agent for detecting mRNA or genomic DNA encoding a polypeptide of the invention is a labeled nucleic acid probe capable of hybridizing to mRNA or genomic DNA encoding a polypeptide of the invention. The nucleic acid probe can be, for example, a full-length cDNA, such as the nucleic acid of SEQ ID NO:1, 2, 19, or 20, or a portion thereof, such as an oligonucleotide of at least 15, 30, 50, 100, 250 or 500 nucleotides in length and sufficient to specifically hybridize under stringent conditions to a mRNA or genomic DNA encoding a polypeptide of the invention. Other suitable probes for use in the diagnostic assays of the invention are described herein.

A preferred agent for detecting a polypeptide of the invention is an antibody capable of binding to a polypeptide of the invention, preferably an antibody with a detectable label. Antibodies can be polyclonal, or more preferably, monoclonal. An intact antibody, or a fragment thereof (e.g., Fab or F(ab')₂) can be used. The term "labeled", with regard to the probe or antibody, is intended to encompass direct labeling of the probe or antibody by coupling (i.e., physically linking) a detectable substance to the probe or antibody, as well as indirect labeling of the probe or antibody by reactivity with another reagent that is directly labeled. Examples of indirect labeling include detection of a primary antibody using a fluorescently labeled secondary antibody and end-labeling of a DNA probe with biotin such that it can be detected with fluorescently labeled streptavidin. The term "biological sample" is intended to include tissues, cells and biological fluids isolated from a subject, as well as tissues, cells and fluids present within a subject. That is, the detection method of the invention can be used to detect mRNA, protein, or genomic DNA in a biological sample *in vitro* as well as *in vivo*. For example, *in vitro* techniques for detection of mRNA include Northern hybridizations and *in situ* hybridizations. *In vitro* techniques for detection of a polypeptide of the invention include enzyme

linked immunosorbent assays (ELISAs), Western blots, immunoprecipitations and immunofluorescence. *In vitro* techniques for detection of genomic DNA include Southern hybridizations. Furthermore, *in vivo* techniques for detection of a polypeptide of the invention include introducing into a subject a labeled antibody directed against the polypeptide. For example, the antibody can be labeled with a radioactive marker whose presence and location in a subject can be detected by standard imaging techniques.

In one embodiment, the biological sample contains protein molecules from the test subject. Alternatively, the biological sample can contain mRNA molecules from the test subject or genomic DNA molecules from the test subject. A preferred biological sample is a peripheral blood leukocyte sample isolated by conventional means from a subject.

In another embodiment, the methods further involve obtaining a control biological sample from a control subject, contacting the control sample with a compound or agent capable of detecting a polypeptide of the invention or mRNA or genomic DNA encoding a polypeptide of the invention, such that the presence of the polypeptide or mRNA or genomic DNA encoding the polypeptide is detected in the biological sample, and comparing the presence of the polypeptide or mRNA or genomic DNA encoding the polypeptide in the control sample with the presence of the polypeptide or mRNA or genomic DNA encoding the polypeptide in the test sample.

The invention also encompasses kits for detecting the presence of a polypeptide or nucleic acid of the invention in a biological sample (a test sample). Such kits can be used to determine if a subject is suffering from or is at increased risk of developing a disorder associated with aberrant expression of a polypeptide of the invention (e.g., a proliferative disorder, e.g., psoriasis or cancer). For example, the kit can comprise a labeled compound or agent capable of detecting the polypeptide or mRNA encoding the polypeptide in a biological sample and means for determining the amount of the polypeptide or mRNA in the sample (e.g., an antibody which binds the polypeptide or an oligonucleotide probe which binds to DNA or mRNA encoding the polypeptide). Kits can also include instructions for observing that the tested subject is suffering from or is at risk of developing a disorder associated with aberrant expression of the polypeptide if the amount of the polypeptide or mRNA encoding the polypeptide is above or below a normal level.

For antibody-based kits, the kit can comprise, for example: (1) a first antibody (e.g., attached to a solid support) which binds to a polypeptide of the invention; and, optionally, (2) a second, different antibody which binds to either the polypeptide or the first antibody and is conjugated to a detectable agent.

For oligonucleotide-based kits, the kit can comprise, for example: (1) an oligonucleotide, e.g., a detectably labeled oligonucleotide, which hybridizes to a nucleic acid sequence encoding a polypeptide of the invention or (2) a pair of primers useful for amplifying a nucleic acid molecule

encoding a polypeptide of the invention. The kit can also comprise, e.g., a buffering agent, a preservative, or a protein stabilizing agent. The kit can also comprise components necessary for detecting the detectable agent (e.g., an enzyme or a substrate). The kit can also contain a control sample or a series of control samples which can be assayed and compared to the test sample contained. Each component of the kit is usually enclosed within an individual container and all of the various containers are within a single package along with instructions for observing whether the tested subject is suffering from or is at risk of developing a disorder associated with aberrant expression of the polypeptide.

2. Prognostic Assays

The methods described herein can furthermore be utilized as diagnostic or prognostic assays to identify subjects having or at risk of developing a disease or disorder associated with aberrant expression or activity of a polypeptide of the invention. For example, the assays described herein, such as the preceding diagnostic assays or the following assays, can be utilized to identify a subject having or at risk of developing a disorder associated with aberrant expression or activity of a polypeptide of the invention, e.g., an immunologic disorder, e.g., asthma, anaphylaxis, or atopic dermatitis. Alternatively, the prognostic assays can be utilized to identify a subject having or at risk for developing such a disease or disorder. Thus, the present invention provides a method in which a test sample is obtained from a subject and a polypeptide or nucleic acid (e.g., mRNA, genomic DNA) of the invention is detected, wherein the presence of the polypeptide or nucleic acid is diagnostic for a subject having or at risk of developing a disease or disorder associated with aberrant expression or activity of the polypeptide. As used herein, a "test sample" refers to a biological sample obtained from a subject of interest. For example, a test sample can be a biological fluid (e.g., serum), cell sample, or tissue.

Furthermore, the prognostic assays described herein can be used to determine whether a subject can be administered an agent (e.g., an agonist, antagonist, peptidomimetic, protein, peptide, nucleic acid, small molecule, or other drug candidate) to treat a disease or disorder associated with aberrant expression or activity of a polypeptide of the invention. For example, such methods can be used to determine whether a subject can be effectively treated with a specific agent or class of agents (e.g., agents of a type which decrease activity of the polypeptide). Thus, the present invention provides methods for determining whether a subject can be effectively treated with an agent for a disorder associated with aberrant expression or activity of a polypeptide of the invention in which a test sample is obtained and the polypeptide or nucleic acid encoding the polypeptide is detected (e.g., wherein the presence of the polypeptide or nucleic acid is diagnostic for a subject that can be

administered the agent to treat a disorder associated with aberrant expression or activity of the polypeptide).

The methods of the invention can also be used to detect genetic lesions or mutations in a gene of the invention, thereby determining if a subject with the lesioned gene is at risk for a disorder characterized aberrant expression or activity of a polypeptide of the invention. In preferred embodiments, the methods include detecting, in a sample of cells from the subject, the presence or absence of a genetic lesion or mutation characterized by at least one of an alteration affecting the integrity of a gene encoding the polypeptide of the invention, or the mis-expression of the gene encoding the polypeptide of the invention. For example, such genetic lesions or mutations can be detected by ascertaining the existence of at least one of: 1) a deletion of one or more nucleotides from the gene; 2) an addition of one or more nucleotides to the gene; 3) a substitution of one or more nucleotides of the gene; 4) a chromosomal rearrangement of the gene; 5) an alteration in the level of a messenger RNA transcript of the gene; 6) an aberrant modification of the gene, such as of the methylation pattern of the genomic DNA; 7) the presence of a non-wild type splicing pattern of a messenger RNA transcript of the gene; 8) a non-wild type level of a the protein encoded by the gene; 9) an allelic loss of the gene; and 10) an inappropriate post-translational modification of the protein encoded by the gene. As described herein, there are a large number of assay techniques known in the art which can be used for detecting lesions in a gene.

In certain embodiments, detection of the lesion involves the use of a probe/primer in a polymerase chain reaction (PCR) (*see, e.g.*, U.S. Patent Nos. 4,683,195 and 4,683,202), such as anchor PCR or RACE PCR, or, alternatively, in a ligation chain reaction (LCR) (*see, e.g.*, Landegran et al. (1988) *Science* 241:1077-1080; and Nakazawa et al. (1994) *Proc. Natl. Acad. Sci. USA* 91:360-364), the latter of which can be particularly useful for detecting point mutations in a gene (*see, e.g.*, Abravaya et al. (1995) *Nucleic Acids Res.* 23:675-682). This method can include the steps of collecting a sample of cells from a patient, isolating nucleic acid (e.g., genomic, mRNA or both) from the cells of the sample, contacting the nucleic acid sample with one or more primers which specifically hybridize to the selected gene under conditions such that hybridization and amplification of the gene (if present) occurs, and detecting the presence or absence of an amplification product, or detecting the size of the amplification product and comparing the length to a control sample. It is anticipated that PCR and/or LCR may be desirable to use as a preliminary amplification step in conjunction with any of the techniques used for detecting mutations described herein.

Alternative amplification methods include: self sustained sequence replication (Guatelli et al. (1990) *Proc. Natl. Acad. Sci. USA* 87:1874-1878), transcriptional amplification system (Kwoh, et al. (1989) *Proc. Natl. Acad. Sci. USA* 86:1173-1177), Q-Beta Replicase (Lizardi et al. (1988) *Bio/Technology* 6:1197), or any other nucleic acid amplification method, followed by the detection of

the amplified molecules using techniques well known to those of skill in the art. These detection schemes are especially useful for the detection of nucleic acid molecules if such molecules are present in very low numbers.

In an alternative embodiment, mutations in a selected gene from a sample cell can be identified by alterations in restriction enzyme cleavage patterns. For example, sample and control DNA is isolated, amplified (optionally), digested with one or more restriction endonucleases, and fragment length sizes are determined by gel electrophoresis and compared. Differences in fragment length sizes between sample and control DNA indicates mutations in the sample DNA. Moreover, the use of sequence specific ribozymes (*see, e.g.*, U.S. Patent No. 5,498,531) can be used to score for the presence of specific mutations by development or loss of a ribozyme cleavage site.

In other embodiments, genetic mutations can be identified by hybridizing a sample and control nucleic acids, e.g., DNA or RNA, to high density arrays containing hundreds or thousands of oligonucleotides probes (Cronin et al. (1996) *Human Mutation* 7:244-255; Kozal et al. (1996) *Nature Medicine* 2:753-759). For example, genetic mutations can be identified in two-dimensional arrays containing light-generated DNA probes as described in Cronin et al., *supra*. Briefly, a first hybridization array of probes can be used to scan through long stretches of DNA in a sample and control to identify base changes between the sequences by making linear arrays of sequential overlapping probes. This step allows the identification of point mutations. This step is followed by a second hybridization array that allows the characterization of specific mutations by using smaller, specialized probe arrays complementary to all variants or mutations detected. Each mutation array is composed of parallel probe sets, one complementary to the wild-type gene and the other complementary to the mutant gene.

In yet another embodiment, any of a variety of sequencing reactions known in the art can be used to directly sequence the selected gene and detect mutations by comparing the sequence of the sample nucleic acids with the corresponding wild-type (control) sequence. Examples of sequencing reactions include those based on techniques developed by Maxim and Gilbert ((1977) *Proc. Natl. Acad. Sci. USA* 74:560) or Sanger ((1977) *Proc. Natl. Acad. Sci. USA* 74:5463). It is also contemplated that any of a variety of automated sequencing procedures can be utilized when performing the diagnostic assays ((1995) *Bio/Techniques* 19:448), including sequencing by mass spectrometry (*see, e.g.*, PCT Publication No. WO 94/16101; Cohen et al. (1996) *Adv. Chromatogr.* 36:127-162; and Griffin et al. (1993) *Appl. Biochem. Biotechnol.* 38:147-159).

Other methods for detecting mutations in a selected gene include methods in which protection from cleavage agents is used to detect mismatched bases in RNA/RNA or RNA/DNA heteroduplexes (Myers et al. (1985) *Science* 230:1242). In general, the technique of Amismatch cleavage entails providing heteroduplexes formed by hybridizing (labeled) RNA or DNA

containing the wild-type sequence with potentially mutant RNA or DNA obtained from a tissue sample. The double-stranded duplexes are treated with an agent which cleaves single-stranded regions of the duplex such as which will exist due to basepair mismatches between the control and sample strands. RNA/DNA duplexes can be treated with RNase to digest mismatched regions, and
5 DNA/DNA hybrids can be treated with S1 nuclease to digest mismatched regions.

In other embodiments, either DNA/DNA or RNA/DNA duplexes can be treated with hydroxylamine or osmium tetroxide and with piperidine in order to digest mismatched regions. After digestion of the mismatched regions, the resulting material is then separated by size on denaturing polyacrylamide gels to determine the site of mutation. *See, e.g.,* Cotton et al. (1988) *Proc. Natl.*
10 *Acad. Sci. USA* 85:4397; Saleeba et al. (1992) *Methods Enzymol.* 217:286-295. In a preferred embodiment, the control DNA or RNA can be labeled for detection.

In still another embodiment, the mismatch cleavage reaction employs one or more proteins that recognize mismatched base pairs in double-stranded DNA (so called ADNA mismatch repair≡ enzymes) in defined systems for detecting and mapping point mutations in cDNAs obtained from
15 samples of cells. For example, the mutY enzyme of *E. coli* cleaves A at G/A mismatches and the thymidine DNA glycosylase from HeLa cells cleaves T at G/T mismatches (Hsu et al. (1994) *Carcinogenesis* 15:1657-1662). According to an exemplary embodiment, a probe based on a selected sequence, e.g., a wild-type sequence, is hybridized to a cDNA or other DNA product from a test cell(s). The duplex is treated with a DNA mismatch repair enzyme, and the cleavage products, if
20 any, can be detected from electrophoresis protocols or the like. *See, e.g.,* U.S. Patent No. 5,459,039.

In other embodiments, alterations in electrophoretic mobility will be used to identify mutations in genes. For example, single strand conformation polymorphism (SSCP) may be used to detect differences in electrophoretic mobility between mutant and wild type nucleic acids (Orita et al. (1989) *Proc. Natl. Acad. Sci. USA* 86:2766; *see also* Cotton (1993) *Mutat. Res.* 285:125-144;
25 Hayashi (1992) *Genet. Anal. Tech. Appl.* 9:73-79). Single-stranded DNA fragments of sample and control nucleic acids will be denatured and allowed to renature. The secondary structure of single-stranded nucleic acids varies according to sequence, and the resulting alteration in electrophoretic mobility enables the detection of even a single base change. The DNA fragments may be labeled or detected with labeled probes. The sensitivity of the assay may be enhanced by using RNA (rather
30 than DNA), in which the secondary structure is more sensitive to a change in sequence. In a preferred embodiment, the subject method utilizes heteroduplex analysis to separate double stranded heteroduplex molecules on the basis of changes in electrophoretic mobility (Keen et al. (1991) *Trends Genet.* 7:5).

In yet another embodiment, the movement of mutant or wild-type fragments in
35 polyacrylamide gels containing a gradient of denaturant is assayed using denaturing gradient gel

electrophoresis (DGGE) (Myers et al. (1985) *Nature* 313:495). When DGGE is used as the method of analysis, DNA will be modified to insure that it does not completely denature, for example by adding a 'GC clamp of approximately 40 bp of high-melting GC-rich DNA by PCR. In a further embodiment, a temperature gradient is used in place of a denaturing gradient to identify differences in the mobility of control and sample DNA (Rosenbaum and Reissner (1987) *Biophys. Chem.* 265:12753).

Examples of other techniques for detecting point mutations include, but are not limited to, selective oligonucleotide hybridization, selective amplification, or selective primer extension. For example, oligonucleotide primers may be prepared in which the known mutation is placed centrally and then hybridized to target DNA under conditions which permit hybridization only if a perfect match is found (Saiki et al. (1986) *Nature* 324:163); Saiki et al. (1989) *Proc. Natl. Acad. Sci. USA* 86:6230). Such allele specific oligonucleotides are hybridized to PCR amplified target DNA or a number of different mutations when the oligonucleotides are attached to the hybridizing membrane and hybridized with labeled target DNA.

Alternatively, allele specific amplification technology which depends on selective PCR amplification may be used in conjunction with the instant invention. Oligonucleotides used as primers for specific amplification may carry the mutation of interest in the center of the molecule (so that amplification depends on differential hybridization) (Gibbs et al. (1989) *Nucleic Acids Res.* 17:2437-2448) or at the extreme 3' end of one primer where, under appropriate conditions, mismatch can prevent or reduce polymerase extension (Prossner (1993) *Tibtech* 11:238). In addition, it may be desirable to introduce a novel restriction site in the region of the mutation to create cleavage-based detection (Gasparini et al. (1992) *Mol. Cell Probes* 6:1). It is anticipated that in certain embodiments amplification may also be performed using Taq ligase for amplification (Barany (1991) *Proc. Natl. Acad. Sci. USA* 88:189). In such cases, ligation will occur only if there is a perfect match at the 3' end of the 5' sequence making it possible to detect the presence of a known mutation at a specific site by looking for the presence or absence of amplification.

The methods described herein may be performed, for example, by utilizing pre-packaged diagnostic kits comprising at least one probe nucleic acid or antibody reagent described herein, which may be conveniently used, e.g., in clinical settings to diagnose patients exhibiting symptoms or family history of a disease or illness involving a gene encoding a polypeptide of the invention. Furthermore, any cell type or tissue, e.g., chondrocytes, in which the polypeptide of the invention is expressed may be utilized in the prognostic assays described herein.

3. Pharmacogenomics

Agents, or modulators which have a stimulatory or inhibitory effect on activity or expression of a polypeptide of the invention as identified by a screening assay described herein can be administered to individuals to treat (prophylactically or therapeutically) disorders associated with aberrant activity of the polypeptide. In conjunction with such treatment, the pharmacogenomics (i.e., the study of the relationship between an individual's genotype and that individual's response to a foreign compound or drug) of the individual may be considered. Differences in metabolism of therapeutics can lead to severe toxicity or therapeutic failure by altering the relation between dose and blood concentration of the pharmacologically active drug. Thus, the pharmacogenomics of the individual permits the selection of effective agents (e.g., drugs) for prophylactic or therapeutic treatments based on a consideration of the individual's genotype. Such pharmacogenomics can further be used to determine appropriate dosages and therapeutic regimens. Accordingly, the activity of a polypeptide of the invention, expression of a nucleic acid of the invention, or mutation content of a gene of the invention in an individual can be determined to thereby select appropriate agent(s) for therapeutic or prophylactic treatment of the individual.

Pharmacogenomics deals with clinically significant hereditary variations in the response to drugs due to altered drug disposition and abnormal action in affected persons. *See, e.g., Linder (1997) Clin. Chem. 43(2):254-266.* In general, two types of pharmacogenetic conditions can be differentiated. Genetic conditions transmitted as a single factor altering the way drugs act on the body are referred to as "altered drug action." Genetic conditions transmitted as single factors altering the way the body acts on drugs are referred to as "altered drug metabolism". These pharmacogenetic conditions can occur either as rare defects or as polymorphisms. For example, glucose-6-phosphate dehydrogenase deficiency (G6PD) is a common inherited enzymopathy in which the main clinical complication is haemolysis after ingestion of oxidant drugs (anti-malarials, sulfonamides, analgesics, nitrofurans) and consumption of fava beans.

As an illustrative embodiment, the activity of drug metabolizing enzymes is a major determinant of both the intensity and duration of drug action. The discovery of genetic polymorphisms of drug metabolizing enzymes (e.g., predicted N-acetyltransferase 2 (NAT 2) and cytochrome P450 enzymes CYP2D6 and CYP2C19) has provided an explanation as to why some patients do not obtain the expected drug effects or show exaggerated drug response and serious toxicity after taking the standard and safe dose of a drug. These polymorphisms are expressed in two phenotypes in the population, the extensive metabolizer (EM) and poor metabolizer (PM). The prevalence of PM is different among different populations. For example, the gene coding for CYP2D6 is highly polymorphic and several mutations have been identified in PM, which all lead to the absence of functional CYP2D6. Poor metabolizers of CYP2D6 and CYP2C19 quite frequently experience exaggerated drug response and side effects when they receive standard doses. If a

metabolite is the active therapeutic moiety, a PM will show no therapeutic response, as demonstrated for the analgesic effect of codeine mediated by its CYP2D6-formed metabolite morphine. The other extreme are the so called ultra-rapid metabolizers who do not respond to standard doses. Recently, the molecular basis of ultra-rapid metabolism has been identified to be due to CYP2D6 gene amplification.

Thus, the activity of a polypeptide of the invention, expression of a nucleic acid encoding the polypeptide, or mutation content of a gene encoding the polypeptide in an individual can be determined to thereby select appropriate agent(s) for therapeutic or prophylactic treatment of the individual. In addition, pharmacogenetic studies can be used to apply genotyping of polymorphic alleles encoding drug-metabolizing enzymes to the identification of an individual's drug responsiveness phenotype. This knowledge, when applied to dosing or drug selection, can avoid adverse reactions or therapeutic failure and thus enhance therapeutic or prophylactic efficiency when treating a subject with a modulator of activity or expression of the polypeptide, such as a modulator identified by one of the exemplary screening assays described herein.

4. Monitoring of Effects During Clinical Trials

Monitoring the influence of agents (e.g., drugs, compounds) on the expression or activity of a polypeptide of the invention (e.g., the ability to modulate aberrant cell proliferation chemotaxis, and/or differentiation) can be applied not only in basic drug screening, but also in clinical trials. For example, the effectiveness of an agent, as determined by a screening assay as described herein, to increase gene expression, protein levels or protein activity, can be monitored in clinical trials of subjects exhibiting decreased gene expression, protein levels, or protein activity. Alternatively, the effectiveness of an agent, as determined by a screening assay, to decrease gene expression, protein levels or protein activity, can be monitored in clinical trials of subjects exhibiting increased gene expression, protein levels, or protein activity. In such clinical trials, expression or activity of a polypeptide of the invention and preferably, that of other polypeptide that have been implicated in for example, a cellular proliferation disorder, can be used as a marker of the immune responsiveness of a particular cell.

For example, and not by way of limitation, genes, including those of the invention, that are modulated in cells by treatment with an agent (e.g., compound, drug or small molecule) which modulates activity or expression of a polypeptide of the invention (e.g., as identified in a screening assay described herein) can be identified. Thus, to study the effect of agents on cellular proliferation disorders, for example, in a clinical trial, cells can be isolated and RNA prepared and analyzed for the levels of expression of a gene of the invention and other genes implicated in the disorder. The levels of gene expression (i.e., a gene expression pattern) can be quantified by Northern blot analysis

or RT-PCR, as described herein, or alternatively by measuring the amount of protein produced, by one of the methods as described herein, or by measuring the levels of activity of a gene of the invention or other genes. In this way, the gene expression pattern can serve as a marker, indicative of the physiological response of the cells to the agent. Accordingly, this response state may be
5 determined before, and at various points during, treatment of the individual with the agent.

In a preferred embodiment, the present invention provides a method for monitoring the effectiveness of treatment of a subject with an agent (e.g., an agonist, antagonist, peptidomimetic, protein, peptide, nucleic acid, small molecule, or other drug candidate identified by the screening assays described herein) comprising the steps of (i) obtaining a pre-administration sample from a
10 subject prior to administration of the agent; (ii) detecting the level of the polypeptide or nucleic acid of the invention in the preadministration sample; (iii) obtaining one or more post-administration samples from the subject; (iv) detecting the level the of the polypeptide or nucleic acid of the invention in the post-administration samples; (v) comparing the level of the polypeptide or nucleic acid of the invention in the pre-administration sample with the level of the polypeptide or nucleic
15 acid of the invention in the post-administration sample or samples; and (vi) altering the administration of the agent to the subject accordingly. For example, increased administration of the agent may be desirable to increase the expression or activity of the polypeptide to higher levels than detected, i.e., to increase the effectiveness of the agent. Alternatively, decreased administration of the agent may be desirable to decrease expression or activity of the polypeptide to lower levels than
20 detected, i.e., to decrease the effectiveness of the agent.

C. Methods of Treatment

The present invention provides for both prophylactic and therapeutic methods of treating a subject at risk of (or susceptible to) a disorder or having a disorder associated with aberrant
25 expression or activity of a polypeptide of the invention. For example, disorders characterized by aberrant expression or activity of the polypeptides of the invention include immunologic disorders. In addition, the nucleic acids, polypeptides, and modulators thereof of the invention can be used to treat immunologic diseases and disorders, including but not limited to inflammatory disorders (e.g., atopic dermatitis). Polypeptides of the invention can also treat diseases associated and bone and
30 cartilage degenerative diseases and disorders (e.g., arthritis, e.g., rheumatoid arthritis), as well as other disorders described herein.

1. Prophylactic Methods

In one aspect, the invention provides a method for preventing in a subject, a disease or
35 condition associated with an aberrant expression or activity of a polypeptide of the invention, by

administering to the subject an agent which modulates expression or at least one activity of the polypeptide. Subjects at risk for a disease which is caused or contributed to by aberrant expression or activity of a polypeptide of the invention can be identified by, for example, any or a combination of diagnostic or prognostic assays as described herein. Administration of a prophylactic agent can occur prior to the manifestation of symptoms characteristic of the aberrancy, such that a disease or disorder is prevented or, alternatively, delayed in its progression. Depending on the type of aberrancy, for example, an agonist or antagonist agent can be used for treating the subject. For example, an antagonist of an A259 protein may be used to treat an arthropathic disorder, e.g., rheumatoid arthritis. The appropriate agent can be determined based on screening assays described herein.

2. Therapeutic Methods

Another aspect of the invention pertains to methods of modulating expression or activity of a polypeptide of the invention for therapeutic purposes. The modulatory method of the invention involves contacting a cell with an agent that modulates one or more of the activities of the polypeptide. An agent that modulates activity can be an agent as described herein, such as a nucleic acid or a protein, a naturally-occurring cognate ligand of the polypeptide, a peptide, a peptidomimetic, or other small molecule. In one embodiment, the agent stimulates one or more of the biological activities of the polypeptide. Examples of such stimulatory agents include the active polypeptide of the invention and a nucleic acid molecule encoding the polypeptide of the invention that has been introduced into the cell. In another embodiment, the agent inhibits one or more of the biological activities of the polypeptide of the invention. Examples of such inhibitory agents include antisense nucleic acid molecules and antibodies. These modulatory methods can be performed *in vitro* (e.g., by culturing the cell with the agent) or, alternatively, *in vivo* (e.g., by administering the agent to a subject). As such, the present invention provides methods of treating an individual afflicted with a disease or disorder characterized by aberrant expression or activity of a polypeptide of the invention. In one embodiment, the method involves administering an agent (e.g., an agent identified by a screening assay described herein), or combination of agents that modulates (e.g., upregulates or downregulates) expression or activity. In another embodiment, the method involves administering a polypeptide of the invention or a nucleic acid molecule of the invention as therapy to compensate for reduced or aberrant expression or activity of the polypeptide.

Stimulation of activity is desirable in situations in which activity or expression is abnormally low or downregulated and/or in which increased activity is likely to have a beneficial effect. Conversely, inhibition of activity is desirable in situations in which activity or expression is

abnormally high or upregulated and/or in which decreased activity is likely to have a beneficial effect.

This invention is further illustrated by the following examples which should not be construed as limiting. The contents of all references, patents and published patent applications cited throughout this application are hereby incorporated by reference.

Deposit of Clones

A clone containing the cDNA molecule encoding human A259, (clone Human I2) was deposited with the American Type Culture Collection, 10801 University Boulevard, Manassas, VA, 20110-2209, on April 2, 1999 as Accession Number 207190, as a single deposit.

A clone containing the cDNA molecule encoding mouse A259, (clone Mouse I2) was deposited with the American Type Culture Collection, 10801 University Boulevard, Manassas, VA, 20110-2209, on April 2, 1999 as Accession Number 207191, as a single deposit.

Equivalents

Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments of the invention described herein. Such equivalents are intended to be encompassed by the following claims.